

FORM PTO-1390 (Modified) (REV 10-95)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER 197317US0PCT
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371			U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR) 09/646897 ✓
INTERNATIONAL APPLICATION NO. PCT/IB99/00941 ✓	INTERNATIONAL FILING DATE 06 April 1999 ✓	PRIORITY DATE CLAIMED 09 April 1998(earliest) ✓	
TITLE OF INVENTION COMPOSITE CO-STRUCTURED OR CO-ADSORBED, MINERAL OR ORGANIC FILLER OR PIGMENT COMPOUNDS AND THE USE THEREOF			
APPLICANT(S) FOR DO/EO/US Patrick A. GANE, et al.			
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:			
<ol style="list-style-type: none"> 1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. 2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. 3. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1). 4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. 5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371 (c) (2)) <ol style="list-style-type: none"> a. <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). b. <input checked="" type="checkbox"/> has been transmitted by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). 6. <input checked="" type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)). 7. <input checked="" type="checkbox"/> A copy of the International Search Report (PCT/ISA/210). 8. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau). b. <input type="checkbox"/> have been transmitted by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input checked="" type="checkbox"/> have not been made and will not be made. 9. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). 10. <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)). 11. <input type="checkbox"/> A copy of the International Preliminary Examination Report (PCT/PEA/409). 12. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)). 			
Items 13 to 18 below concern document(s) or information included:			
<ol style="list-style-type: none"> 13. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. 14. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. 15. <input checked="" type="checkbox"/> A FIRST preliminary amendment. A SECOND or SUBSEQUENT preliminary amendment. 16. <input type="checkbox"/> A substitute specification. 17. <input type="checkbox"/> A change of power of attorney and/or address letter. 18. <input type="checkbox"/> Certificate of Mailing by Express Mail 19. <input checked="" type="checkbox"/> Other items or information: 			
Request for Consideration of Documents Cited in International Search Report Notice of Priority Drawings (5 Sheets)			

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 09/646897)	INTERNATIONAL APPLICATION NO. PCT/IB99/00941	ATTORNEY'S DOCKET NUMBER 197317US0PCT
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20. The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)):

- | | |
|--|-----------|
| <input checked="" type="checkbox"/> Search Report has been prepared by the EPO or JPO | \$860.00 |
| <input checked="" type="checkbox"/> International preliminary examination fee paid to USPTO (37 CFR 1.482) | \$690.00 |
| <input type="checkbox"/> No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)) | \$710.00 |
| <input type="checkbox"/> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO | \$1000.00 |
| <input type="checkbox"/> International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4) | \$100.00 |

ENTER APPROPRIATE BASIC FEE AMOUNT =

\$860.00

Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☒ 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).

\$130.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	
Total claims	37 - 20 =	17	x \$18.00	\$306.00
Independent claims	1 - 3 =	0	x \$80.00	\$0.00
Multiple Dependent Claims (check if applicable). <input type="checkbox"/>				\$0.00

TOTAL OF ABOVE CALCULATIONS =

\$1,296.00

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) (check if applicable). ☐

\$0.00

SUBTOTAL =

\$1,296.00

Processing fee of \$130.00 for furnishing the English translation later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).

\$0.00

TOTAL NATIONAL FEE =

\$1,296.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). ☐

\$0.00

TOTAL FEES ENCLOSED =

\$1,296.00

Amount to be:	\$
refunded	
charged	\$

☒ A check in the amount of \$1,296.00 to cover the above fees is enclosed.

☐ Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.

☒ The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. **15-0030** A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:



22850

Surinder Sachar
Registration No. 34,423

Surinder Sachar
SIGNATURE

Norman F. Oblon
NAME

24,618
REGISTRATION NUMBER

10-10-00
DATE

09/646897

422 Rec'd PCT/PTO 1 0 OCT 2000

197317US0 PCT

IN THE UNITED STATES PATENT & TRADEMARK OFFICE

IN RE APPLICATION OF :
PATRICK A. GANE ET AL : ATTN: APPLICATION DIVISION
SERIAL NO: NEW PCT APPLICATION :
(Based on PCT/IB99/00941)
FILED: HEREWITH :
FOR: COMPOSITE CO-STRUCTURED :
OR CO-ADSORBED, MINERAL OR
ORGANIC FILLER OR PIGMENT
COMPOUNDS AND THE USE :
THEREOF

PRELIMINARY AMENDMENT

ASSISTANT COMMISSIONER FOR PATENTS
WASHINGTON, D.C. 20231

SIR:

Prior to examination on the merits, please amend the above-identified application as follows.

IN THE TITLE

Please replace the title with the following substitute title:

— COMPOSITE COMPOSITIONS OF CO-STRUCTURED OR CO-ADSORBED
ORGANIC OR MINERAL PIGMENTS OR FILLERS AND THEIR USES--

IN THE CLAIMS

Please amend the claims as follows:

Claim 5, lines 1-2, replace "one of claims 1 to 4" with --claim 1--.

Claim 6, lines 1-2, replace "any one of claims 1 to 5" with --claim 1--.

Claim 7, lines 1-2, replace "any one of claims 1 to 6" with --claim 1--.

--8. (Amended) A composite as claimed in claim 1 [any one of claims 1 to 6], characterized in that the binding agent is selected from among the acrylic or vinyl polymers and/or copolymers [...] in their totally acid state or partially neutralised, or totally neutralised, obtained by polymerisation, in the acid state in the presence of one at least of the mineral or organic particles of the composite compound and optionally in the presence of [the] a binding agent [as claimed in claim 7,] ~~selected from the group consisting of~~ at least one of the monomers such as acrylic acid and/or methacrylic acid, itaconic, crotonic, fumaric acid, maleic anhydride, or isocrotonic, aconitic, mesaconic, sinapic, undecylenic, angelic acid and/or their respective esters, acrylamido methyl propane sulphonic acid, acrolein, acrylamide and/or methacrylamide, methacrylamido propyl trimethyl ammonium chloride or sulphate, methacrylate of trimethyl ammonium ethyl chloride or sulphate, as well as their acrylate and acrylamide counterparts, quaternised or not, and/or dimethyldiallyl chloride, vinylpyrrolidone or such as the unsaturated linear or branched fatty acids or the linear or branched fatty alcohols or the unsaturated linear or branched or cyclic fatty amines, or the quaternary salts, preferably with unsaturated linear or branched fatty chains or vegetable origin or not.

Claim 9, lines 1-2, replace "any one of claims 1 to 8" with --claim 1--.

Claim 10, lines 1-2, replace "any one of claims 1 to 8" with --claim 1--.

Claim 11, lines 1-2, replace "any one of claims 1 to 10" with --claim 1--.

Claim 12, lines 1-2, replace "any one of claims 1 to 11" with --claim 1--.

Claim 13, lines 1-2, replace "any one of claims 1 to 12" with --claim 1--.

Claim 14, lines 1-2, replace "any one of claims 1 to 12" with --claim 1--.

Claim 15, lines 1-2, replace "any one of claims 1 to 14" with --claim 1--.

- ° Claim 16, lines 1-2, replace "any one of claims 1 to 14" with --claim 1--.
- ° Claim 17, lines 1-2, replace "any one of claims 1 to 14" with --claim 1--.
- Claim 18, lines 3-4, replace "any one of claims 1 to 14" with --claim 1--.
- Claim 21, lines 2-3, replace "any one of claims 1 to 14" with --claim 1--.
- Claim 24, lines 1-2, replace "any one of claims 21 to 23" with --claim 11--.
- Claim 25, lines 1-2, replace "any one of claims 21 to 23" with --claim 11--.
- Claim 26, lines 1-2, replace "any one of claims 21 to 23" with --claim 21--.
- Claim 27, lines 1-2, replace "any one of claims 21 to 23" with --claim 21--.
- Claim 28, lines 1-2, replace "any one of claims 21 to 23" with --claim 21--.
- Claim 29, lines 5-6, replace "any one of claims 1 to 14" with --claim 1--.
- Claim 33, line 2, replace "any one of claims 29 to 31" with --claim 29--.
- Claim 34, lines 2-3, replace "any one of claims 1 to 14" with --claim 1--.

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COMPOSITE CO-STRUCTURED OR CO-ADSORBED, MINERAL OR
ORGANIC FILLER OR PIGMENT COMPOUNDS AND THE USE THEREOF

5 The present invention relates to the field of composite
compounds of mineral or organic fillers or pigments
containing at least two mineral or organic fillers or
pigments of different natures and the uses thereof in the
paper industry for manufacturing paper, for filling or
10 coating purposes, or for any other surface treatment of
the paper, as well as in the field of aqueous and non-
aqueous paints and in the plastics industry.

Composite pigments or fillers are commonly used these
15 days for the manufacture of all types of paper, filling
and coating or any other surface treatment of the paper
in order to improve the quality of the paper in terms of
its properties of opacity, whiteness and brightness of
the sheets of paper, for example, or to improve the
20 printability characteristics.

One widely used technique of producing composite pigments
or fillers consists in mixing a mineral filler, such as a
natural calcium carbonate for example, with a mineral
25 filler such as talc (FR 2 526 061) or alternatively a
mineral filler such as talc with another mineral filler
such as calcined kaolin (EP 0 365 502).

Two other types of process are also known from the prior
30 art for producing composite pigments or fillers meeting
the requisite criteria for use in the paper industry.

A first category of these processes known from the prior
art involves forming networks between the pigment
35 particles, thereby creating numerous internal voids which

then enhance the optical properties of the pigment fillers, often measured by a light scattering coefficient S.

5 Accordingly, patent WO 92/08755 describes a method of forming aggregates by flocculation and optionally by in situ precipitation of calcium carbonate, this flocculation occurring consecutively with the ionic interactions produced by using anionic polymers with a high molecular weight to flocculate the mineral particles to which multivalent cations such as the calcium ion are added on the surface.

Similarly, patent US 5.449.402 discloses a product obtained by a method of creating internal voids based on ionic or electrostatic interactions, whilst US 5.454.864 or US 5.344.487 or EP 0 573 150 propose a composite pigment, the preparation of which is based on the attraction forces of ions.

These methods based on ionic attraction forces are sensitive to the ionic forces at play in the formulas used for paper coating colors or for paper filling and there is no guarantee that these pigments can be used in applications such as paper coating or paper filling.

A second category of these methods known from the prior art as a means of producing pigments with improved optical characteristics is based on using organic compounds of silicon (US 4.818.294; US 5.458.680) or chloride-based compounds (US 4.820.554; US 4.826.536; WO 97/24406).

Finally, a last known method of improving whiteness (WO 97/32934) consists in coating the pigment particles with

another pigment particle such as very fine particles of precipitated calcium carbonate. However, a method of this type is not based on using an organic binding agent which creates a co-structure.

5 Faced with this problem of improving optical properties, such as opacity, whiteness, colouring or brightness, for example, or improving printability characteristics, the applicant has developed, for the purposes of this
10 invention, composite compounds, which are dry or non-aqueous or aqueous, of mineral or organic fillers or pigments which will improve at least one of the optical properties or printability required in the various fields of application, whilst providing a macroscopically
15 homogeneous and stable compound in spite of the ionic forces present in the known formulas, such as offset or rotogravure paper coating colors or paper filling.

20 Accordingly, one of the objectives of the invention is to provide dry or non-aqueous or aqueous composite compounds of mineral or organic fillers or pigments containing at least two mineral or organic fillers or pigments of a physically or chemically different nature.

25 The composite compounds of organic or mineral fillers or pigments proposed by the invention and constituting one of the objects thereof, which exhibit the qualities outlined above, are characterised in that they contain

30 a) at least two organic or mineral fillers or pigments, at least one of which has a surface having at least one hydrophilic site and the other at least having a surface with at least one organophilic site

35 b) at least one binding agent

and in that they are co-structured or co-adsorbed, i.e. the different mineral or organic particles exhibit a structural cohesion imparted by creating a bond or an adhesion between at least two particles with different surface states.

Consequently, throughout this description the words co-structure or co-structured or co-adsorbed are used by the applicant as meaning the creation of a bond between at least two fillers or pigments of any type by forming a structure comparable to a bond or an adhesion between the surface of a filler or pigment having at least one hydrophilic site and the surface of the other filler or pigment having at least one organophilic site using a binding agent which is an organic compound. This binder may be supported by a gas such as air or any other gas.

Furthermore, another objective of the invention is to develop stable composite compounds which can be transported and stored for several weeks.

Yet another objective of the invention is to develop a composite compound of a macroscopically homogeneous structure, this being achieved by macroscopically stable paper coating colors which contain the above-mentioned composite aqueous compounds.

Moreover, another objective of the invention is the use of these composite compounds as organic or mineral fillers or pigments for the manufacture of paper, filling and/or coating and/or any other compound used to treat the surface of paper, as well as in the field of paints and the field of plastics.

Finally, another objective of the invention is to provide suspensions containing the composite compounds proposed by the invention and to provide paper coating colors or compounds for treating the surface of paper or uncoated filling compounds that will improve at least one of the properties, such as opacity, whiteness, brightness or printability.

It should be pointed out that the improvement made to these properties will depend on the field in which the skilled person wishes to adapt the properties to suit the desired application.

These objectives are achieved by placing the surface of one of the pigments or fillers in contact with the surface of the other of the pigments or fillers in the presence of a binding agent so that this contact causes a structure to form, between at least two mineral or organic particles of a different physical or chemical nature, i.e. causes a structure to form between them, at least one of them having a surface with at least one hydrophilic site and at least one of them having a surface with at least one organophilic site.

By mineral or organic particle having a surface with at least one hydrophilic site, the applicant means a mineral or organic particle which is partially or totally wettable by polar substances without the influence of any external component and more specifically is partially or totally wettable by water.

The mineral or organic particles having a surface with at least one hydrophilic site may be of very different physical or chemical natures, such as natural calcium carbonate, for example chalk, calcite, marble or any

other form of natural calcium carbonate, which may be obtained from a recycling process in particular, precipitated calcium carbonate, the dolomites, crystalline or amorphous aluminium hydroxides, natural or synthetic precipitated silicates, calcium sulphate, titanium dioxide, satin white, the wollastonites, huntite, calcined clays derived from recycling for example, starch or any other type of organophilic organic or mineral particles that have undergone physical processing such as Corona or chemical processing in order to produce at least one hydrophilic site.

By mineral or organic particle having a surface with at least one organophilic site, the applicant means a mineral or organic particle which is partially or totally wettable by an organic fluid or an organic substance, this wettability being independent of adsorption mechanisms such as electrostatic attraction or sequestration.

By mineral or organic particle having a surface with at least one organophilic site, the applicant means in particular a mineral or organic particle of very different physical or chemical natures such as talcs, micas, calcined clays or not, or zinc oxide or transparent iron pigments or colouring pigments such as phthalocyanine blue, polystyrene-based synthetic pigments, urea-formol resins, carbon black, cellulose fibres and flours or any other type of hydrophilic mineral or organic particles but which, after chemical or physical processing, have at least one organophilic site, i.e. are wettable by an organic fluid or an organic substance.

It should be pointed out that the quantities and ratios

by dry weight of the various fillers or pigments making up the composite compounds proposed by the invention vary from 0.1% to 99.9% depending on the nature of the different pigments or fillers, and preferably from 25% to 95% by dry weight, relative to the total dry weight of the fillers or pigments, of mineral or organic fillers or pigments having a surface with at least one hydrophilic site and preferably between 75% and 5% by dry weight, relative to the total dry weight of the fillers or pigments, of mineral or organic fillers or pigments having a surface with at least one organophilic site.

The creation of a bond or a co-structure can be seen in the rheological behaviour of the composite compounds and by the properties of homogeneity in paper coating colors or in the printability of the paper.

It also increases the opacity of sheets of wood free paper filled at a rate of 75.5 g/m² with the composite compounds of the invention. This opacity is measured in compliance with the DIN 53146 standard using an Elrepho 2000 spectrophotometer made by Datacolor AG (Switzerland).

Accordingly, the composite compounds of mineral or organic fillers or pigments proposed by the invention are characterised in that they are co-structured or co-adsorbed, i.e. they have a high yield stress, as determined by a Stress Tech[®] machine, namely higher than and preferably at least four times higher than that of the standard mixture of corresponding fillers or pigments.

The composite compounds of mineral or organic fillers or pigments proposed by the invention are also characterised in that the different mineral or organic particles exhibit a cohesion which reflects the macroscopic homogeneity of the suspension of the composite compound and/or coating color containing the composite compound. This macroscopic homogeneity is expressed by measuring the content of one of the pigments or fillers at two different points of the suspension or of the coating color after it has been left to rest for several hours or several days.

Furthermore, the composite compounds of mineral or organic fillers or pigments proposed by the invention are characterised in that they contain at least one binding agent. This binding agent is an organic compound, which might be supported by a gas such as air or any other gas. This binding agent, an organic compound, must be partially or totally wettable by the surfaces of the pigments or fillers with which it is placed in contact. By preference, this binding agent is selected from among the acrylic or vinyl polymers and/or copolymers or polycondensates or polyaddition products such as the polymers or copolymers for example, in their totally acid state or partially neutralised or totally neutralised by neutralising agents containing monovalent or polyvalent cations or mixtures thereof, by one at least of the monomers such as acrylic and/or methacrylic, itaconic, crotonic, fumaric acid, maleic anhydride or isocrotonic, aconitic, mesaconic, sinapic, undecylenic, angelic acid and/or the respective esters thereof, acrylamido methyl propane sulphonic acid, acrolein, acrylamide and/or methacrylamide, methacrylamido propyl trimethyl ammonium chloride or sulphate, methacrylate of trimethyl ammonium ethyl chloride or sulphate, as well as their acrylate and

acrylamide counterparts, quaternised or not, and/or dimethyldiallyl chloride, vinylpyrrolidone or a binding agent selected from among the linear or branched fatty acids or the linear or branched fatty alcohols or the linear or branched or cyclic fatty amines, saturated or not, or selected from among the quaternary salts preferably with linear or branched fatty chains of vegetable origin or not.

This binding agent may also be selected from among at least one of the above-mentioned monomers or mixtures thereof in the form of the monomer or monomers itself/themselves, polymerised in the presence of at least one of the mineral or organic particles.

Furthermore, it should be noted that optimisation of the molecular weight of the binding agent will depend on its chemical nature.

For the purposes of the invention, the quantity of binding agent used in the composite compound is from 0.01% to 10%, preferably from 0.10% to 1.5% by dry weight relative to the total dry weight of the fillers or pigments.

The composite compounds proposed by the invention may optionally be dispersed in water, in water-solvent mixtures or in other solvents using one or more dispersing agents known to the person skilled in the art, amongst others those described in patents EP 0 100 947, EP 0 542 643 or EP 0 542 644.

It is also important to note that the co-structured composite compounds proposed by the invention are compatible with other aqueous mineral or organic filler

compounds, i.e. they form a stable and homogeneous mixture when they are simply mixed with these other suspensions, whereas it would be impossible to produce a homogeneous suspension if a co-structured composite compound as proposed by the invention were not used.

The paper coating colors and/or paper surface treatment compounds as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous or non-aqueous paint compositions proposed by the invention are prepared in a manner known to the person skilled in the art by mixing in water the dry or non-aqueous or aqueous mineral or organic composite compounds of fillers or pigments proposed by the invention and one or more binders of natural or synthetic origin such as starch, carboxymethyl cellulose, the polyvinyl alcohols, for example, or latex or polymeric dispersions of the styrene-butadiene or styrene-acrylate type or acrylic or vinyl or other polymeric dispersions.

In a known manner, the paper coating colors and/or paper surface treatment compounds as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous or non-aqueous paint compositions may also contain the usual additives such as rheology modifiers, organic fillers, anti-foaming agents, optical brighteners, biocide agents, lubricants, alkaline hydroxides, colourings and others.

In addition, the aqueous suspensions containing the composite compounds, the paper coating colors and/or the paper surface-treatment compounds as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous or non-aqueous paint compositions or the uncoated filling compositions proposed by the invention

are characterised in that they contain dry or non-aqueous or aqueous composite compounds as proposed by the invention.

5 The aqueous suspensions containing the composite compounds, the paper coating colors and/or the paper surface-treatment compounds as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous paint or non-aqueous compositions as proposed by
10 the invention are also characterised in that they are macroscopically homogeneous.

This macroscopic homogeneity is determined by measuring the quantity of one of the fillers at the surface and the
15 base of the flask containing the coating color diluted 40% or 20% with dry substance.

A comparison of this quantity of one of the fillers at the two "top" and "bottom" points of the compound
20 proposed by the invention with the quantity of one of the fillers at these two "top" and "bottom" points of the standard mixture will demonstrate the fact that there is virtually no migration of one of the fillers to a preferred part of the compound of the invention, contrary
25 to the situation that would arise with a standard mixture.

This macroscopic homogeneity of the composite compounds of the invention produces improved homogeneity in the
30 sheet of paper due to a more even retention and distribution.

Moreover, the aqueous suspensions containing the composite compounds of the invention, the paper coating
35 colors of the invention or the paper surface-treatment

compounds of the invention are characterised in that the yield stress, as determined by a Stress Tech[®] machine, is higher than and preferably at least four times higher than the corresponding standard mixtures of fillers or pigments.

Furthermore, the aqueous suspensions of the invention or the coating colors of the invention or the paper surface-treatment compounds of the invention or the uncoated filling compounds of the invention will also exhibit an improvement in at least one of the optical properties such as opacity or whiteness or brightness or in the printability or print density properties.

Likewise, the aqueous or non-aqueous paint compositions containing the composite compounds of the invention have the advantage of an increased opacity.

Accordingly, by preference, the aqueous suspensions containing the composite compounds of the invention or the coating colors of the invention or the paper surface-treatment compounds of the invention as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous or non-aqueous paint compositions are characterised in that they have a higher light scattering coefficient S than that of the corresponding standard mixtures.

The uncoated filling compounds proposed by the invention are preferably characterised in that they exhibit a higher opacity, determined in compliance with the DIN 53416 standard, than that of the corresponding standard mixtures.

Similarly, by preference, the aqueous suspensions containing the composite compounds of the invention or the coating colors of the invention or the paper surface-treatment compound of the invention or the uncoated filling compounds of the invention are characterised in that they have a higher whiteness, determined in compliance with the TAPPI T452 ISO 2470 standard, than that of the corresponding standard mixtures.

By preference, the coating colors or paper surface-treatment compounds proposed by the invention are characterised in that they exhibit a higher TAPPI 75° brightness according to Lehmann than that of a coating color containing standard suspensions of corresponding mixtures.

Finally, and by preference, the coating colors or paper surface-treatment compounds or uncoated filling compounds proposed by the invention are characterised in that the curve, plotted on the basis of the ISIT printability test, the conducting of which will be explained in example 9, representing the ink tack force as a function of time, exhibits less marked rising and falling slopes and a higher maximum value than coating colors or paper surface-treatment compounds or uncoated filling compounds containing standard suspensions of corresponding mixtures.

Furthermore, sheets of paper which contain in the mass the composite compounds proposed by the invention are characterised in that they exhibit a higher whiteness, determined in compliance with the TAPPI T452 ISO 2470 standard, than sheets of paper containing in the mass standard suspensions of mixtures of corresponding fillers or pigments and in that they exhibit a higher opacity,

measured in compliance with the DIN 53146 standard, than sheets of paper containing standard suspensions of mixtures of corresponding fillers or pigments.

- 5 The scope and interest of the invention will be more readily understood from the examples set out below, which are not intended to be restrictive, particularly in terms of the order in which the various constituents of the composite compounds are added.

10

EXAMPLE 1 :

This example relates to the preparation of composite compounds containing different pigments or fillers.

15

The specific viscosities mentioned in all the examples are determined by the method defined in EP 0 542 643.

20

Test N° 1 :

25

This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 72% aqueous suspension of a Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μm as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 250 grams dry weight of an aqueous suspension of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100 containing 0.08% by dry weight of soda, 1.4% by dry weight of an alkylene polyoxide and 0.15% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 in order to produce an aqueous suspension with a 70% concentration of a marble-talc mixture.

30

Test N° 2 :

For this test, illustrating the invention, the co-structured aqueous compound of the invention is prepared by adding in a mixer and under agitation:

- 5 - 750 grams dry of Norwegian marble having a grain size such that 75% of the particles are of a diameter of less than $1\mu\text{m}$ as measured on the Sedigraph 5100,
- 250 grams dry weight of a talc from Finland with a grain size such that 45% of the particles are of a diameter of less than $2\mu\text{m}$ as measured on the Sedigraph 5100,
- 10 - 5 grams dry weight of an acrylic copolymer binding agent of a monomer composition comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide,
- 15 - the quantity of water needed to make up the co-structured aqueous compound of the invention to a 65% concentration of dry substance.

20 After agitating for 30 minutes and after formation of the co-structure between the grains of marble and talc with the aid of the binding agent, 5.2 grams dry weight of a dispersing agent of the prior art, namely a polyacrylate partially neutralised with soda and having a specific
25 viscosity of 0.5, are added to the compound of the invention, the balance being added in the form of soda and water necessary to produce an aqueous suspension of the composite compound of the invention with a concentration of dry substance equal to 59.1% with a pH
30 of between 9 and 10.

Test N° 3 :

This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 72% aqueous

suspension of Champagne chalk with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μm as measured on the Sedigraph 5100 and containing 0.80% by dry weight of a sodium polyacrylate with a specific viscosity equal to 0.5 with 250 grams dry weight of an aqueous suspension of a Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100 containing 0.08% by dry weight of soda, 1.4% by dry weight of an alkylene polyoxide and 0.15% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 in order to produce an aqueous suspension with a 62.1% concentration of a chalk-talc mixture.

Test N° 4 :

This test, illustrating the invention, is conducted in the same manner and using the same equipment as test N° 2 replacing the marble with Champagne chalk of the same grain size.

This produces an aqueous suspension of the co-structured composite compound of the invention (75% by dry weight of chalk - 25% by dry weight of talc) having a 57% concentration of dry substance.

Test N° 5 :

This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 51% aqueous suspension of precipitated calcium carbonate with a grain size equivalent to one in which 60% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100 and dispersed with 0.3% by dry weight of a sodium polyacrylate with a specific viscosity equal to 0.7 with 250 grams dry weight of an aqueous suspension of

a Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μ m as measured on the Sedigraph 5100 containing 0.08% by dry weight of soda, 1.4% by dry weight of an alkylene polyoxide and 0.15% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 in order to produce an aqueous suspension with a 54.5% concentration of a precipitated calcium carbonate-talc mixture.

10 Test N° 6 :

This test, illustrating the invention, is conducted in the same manner and using the same equipment as test N° 2 replacing the marble with precipitated calcium carbonate with a grain size equivalent to one in which 60% of the particles are of a diameter of less than 2 μ m.

This produces an aqueous suspension of the co-structured composite compound of the invention (75% by dry weight of precipitated calcium carbonate - 25% by dry weight of talc) having a 58% concentration of dry substance.

20 Test N° 7 :

This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μ m as measured on the Sedigraph 5100 and containing 1.00% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 250 grams dry weight of an aqueous suspension of an Austrian mica with a grain size equivalent to one in which 18% of the particles are of a diameter of less than 1 μ m as measured on the Sedigraph 5100 containing 0.25% by dry weight of a sodium polyacrylate with a specific viscosity

of 0.4 in order to produce an aqueous suspension with a 68.6% concentration of a marble-mica mixture.

Test N° 8 :

- 5 This test, illustrating the invention, is conducted in the same manner and using the same equipment as test N° 2 replacing the talc with Austrian mica with a grain size equivalent to one in which 18% of the particles are of a diameter of less than $1\mu\text{m}$.
- 10 This produces an aqueous suspension of the co-structured composite compound of the invention (75% by dry weight of marble - 25% by dry weight of mica) having a 61.3% concentration of dry substance.

Test N° 9 :

- 15 This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than $1\mu\text{m}$ as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 250 grams dry weight of an aqueous suspension of an English kaolin with a grain size equivalent to one in which 64%
- 20 of the particles are of a diameter of less than $1\mu\text{m}$ as measured on the Sedigraph 5100 containing 0.2% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 in order to produce an aqueous suspension with a 70.2% concentration of a marble-kaolin mixture.

Test N° 10 :

30 This test, illustrating the invention, is conducted in the same manner and using the same equipment as test N° 2

replacing the talc with English kaolin having a grain size equivalent to one in which 64% of the particles are of a diameter of less than $1\mu\text{m}$ as measured by the Sedigraph 5100.

- 5 This produces an aqueous suspension of the co-structured composite compound of the invention (75% by dry weight of marble - 25% by dry weight of kaolin) having a 62.1% concentration of dry substance.

10 Test N° 11 :

- This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than $1\mu\text{m}$ as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 250 grams dry weight of an aqueous suspension of a titanium dioxide of the rutile type with a grain size equivalent to one in which 86% of the particles are of a diameter of less than $1\mu\text{m}$ as measured on the Sedigraph 5100 containing 0.32% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 in order to produce an aqueous suspension with a 71.5% concentration of a marble-titanium dioxide mixture.
- 25

Test N° 12 :

- For this test, illustrating the invention, the co-structured compound is prepared in the same manner and using the same equipment as test N° 2 replacing the talc with titanium dioxide of the rutile type having a grain size equivalent to one in which 86% of the particles are of a diameter of less than $1\mu\text{m}$.
- 30

After the co-structure has formed between the grains of marble and titanium dioxide with the aid of a binding agent, 0.15% by dry weight of a dispersing agent known from the prior art is added, namely a sodium polyacrylate with a specific viscosity of 0.5.

This produces an aqueous suspension of the co-structured composite compound of the invention (75% by dry weight of marble - 25% by dry weight of titanium dioxide) having a 58.8% concentration of dry substance.

Test N°13 :

This test, illustrating the prior art, is a standard mixture of 750 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μm as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with :

- 125 grams dry weight of an aqueous suspension of English kaolin with a grain size equivalent to one in which 64% of the particles are of a diameter of less than 1 μm as measured on the Sedigraph 5100 and containing 0.3% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4

- 125 grams dry weight of an aqueous suspension of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100 and containing 0.08% by dry weight of soda, 1.4% by dry weight of an alkylene polyoxide and 0.15% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 in order to produce an aqueous suspension with a 70.2% concentration of dry substance in a marble-kaolin-talc mixture.

Test N° 14 :

This test, illustrating the invention, is conducted in the same manner and using the same equipment as test N° 2 replacing half of the talc with English kaolin having a grain size equivalent to one in which 64% of the particles are of a diameter of less than 1 μ m as measured by the Sedigraph 5100.

This produces an aqueous suspension of the co-structured composite compound of the invention (75% by dry weight of marble - 12.5% by dry weight of kaolin - 12.5% by dry weight of talc) having a 60.0% concentration of dry substance.

Test N° 15 :

This test, illustrating the prior art, is a standard mixture of 800 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μ m as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 200 grams dry weight of an aqueous suspension of crystalline aluminium hydroxide with a grain size equivalent to one in which 72% of the particles are of a diameter of less than 2 μ m as measured on the Sedigraph 5100 and containing 0.3% by dry weight of a sodium polyacrylate with a specific viscosity of 0.7 in order to produce an aqueous suspension with a 70.9% concentration of dry substance in a marble-aluminium hydroxide mixture.

Test N° 16 :

For this test, illustrating the invention, the co-

structured aqueous compound proposed by the invention is prepared by introducing into a mixer under agitation:

- 800 grams dry weight of a Norwegian marble with a grain size such that 75% of the particles are of a diameter of less than 1 μm as measured on the Sedigraph 5100,
- 200 grams dry weight of crystalline aluminium hydroxide with a grain size such that 72% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph,
- 4 grams dry weight of an acrylic copolymer binding agent of a monomer composition comprising 90% by dry weight of acrylic acid and 10% by dry weight of triethylphenol methacrylate with 25 moles of ethylene oxide,
- the quantity is made up with the water necessary to produce a co-structured aqueous compound of the invention with a 65% concentration of dry substance.

After 30 minutes of agitation and once the co-structure has formed between the grains of marble and aluminium hydroxide with the aid of the binding agent, 5.6 grams by dry weight of a dispersing agent known from the prior art are added to the compound of the invention, namely a polyacrylate partially neutralised with soda and of a specific viscosity of 0.5, as well as the soda and water required to obtain an aqueous suspension of the composite compound of the invention having a concentration of dry substance equal to 60.3% and with a pH value of between 9 and 10.

Test N° 17 :

This test, illustrating the prior art, is a standard mixture of 800 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a

diameter of less than 1 μm as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 200 grams dry weight of an aqueous suspension of a urea-formol condensate with a specific surface area equal to 17 m^2/g as measured by the BET method (DIN 66132) and containing 0.5% by dry weight of a sodium polyacrylate with a specific viscosity of 0.7 in order to produce an aqueous suspension with a 45.1% concentration of dry substance in a marble-urea-formol condensate mixture.

Test N° 18 :

This test, illustrating the invention, is conducted in the same manner and using the same equipment in all respects as test N° 16 replacing the aluminium hydroxide with a urea-formol condensate having a specific surface area of 17 m^2/g as measured by the BET method (DIN 66132).

This produces an aqueous suspension of the co-structured composite compound of the invention (80% by dry weight of marble - 20% by dry weight of a urea-formol condensate) having a 51.2% concentration of dry substance.

Test N° 19 :

This test, illustrating the prior art, is a standard mixture of 800 grams dry weight of a 72% aqueous suspension of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μm as measured on the Sedigraph 5100 and containing 1% by dry weight of an acrylic copolymer with a specific viscosity equal to 0.8 with 200 grams dry weight of an aqueous suspension of bleached cellulose with a grain size equivalent to one in which

99% of the particles are of a diameter of less than 75 μm as measured by an airflow sieve of the Alpine LS 200 type and containing 0.5% by dry weight of a sodium polyacrylate with a specific viscosity of 0.7 in order to produce an aqueous suspension with a 44.8% concentration of dry substance in a marble-bleached cellulose mixture.

Test N° 20 :

10 This test, illustrating the invention, is conducted in the same manner and using the same equipment in all respects as test N° 16 replacing the aluminium hydroxide with bleached cellulose with a grain size equivalent to on in which 99% of the particles are of a diameter of less than 75 μm as measured by an airflow sieve of the Alpine LS 200 type.

15 This produces an aqueous suspension of the co-structured composite compound of the invention (80% by dry weight of marble - 20% by dry weight of bleached cellulose) having a 46.9% concentration of dry substance.

Test N° 21 :

25 This test, illustrating the prior art, is a standard mixture of 500 grams dry weight of a Champagne chalk with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100 with 500 grams of an Australian talc with a grain size equivalent to one in which 25% of the particles are of a diameter of less than 2 μm as measured by the Sedigraph 5100 to produce a powdered chalk-talc mixture with a 100% concentration of dry substance.

Test N°22 :

For this test, illustrating the invention, the co-structured compound is prepared in a powdered form as proposed by the invention, by introducing into a mixer under agitation:

- 500 grams dry weight of a Champagne chalk of a grain size such that 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100,
- 500 grams dry weight of a talc from Australia of a grain size such that 25% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100,
- 10 grams dry weight of an acrylic copolymer binding agent of a monomer composition comprising 90% by dry weight of acrylic acid and 10% by dry weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide.

Test N°23 :

This test, illustrating the prior art, is a simple mixture of 900 grams dry weight of an aqueous suspension of a Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph 5100 and containing 0.08% by dry weight of soda, 1.4% by dry weight of an alkylene polyoxide and 0.15% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4 with 100 grams dry weight of an American kaolin with a grain size equivalent to one in which 91% of the particles are of a diameter of less than 0.5 μm as measured by the Sedigraph 5100 to produce an aqueous suspension with a 67.8% concentration of a talc-kaolin mixture.

EXAMPLE 2 :

This example illustrates the preparation of composite compounds as proposed by the invention using various ratios of pigments or fillers.

To this end, the composite compounds proposed by the invention are prepared using the same method and the same equipment as in test N° 2 with the exception of the quantity of water, which is added all at once, to produce the final concentration of dry substance and comprising:

Test N° 24 :

95% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 62% of the particles are of a diameter of less than $1\mu\text{m}$ as measured with the Sedigraph 5100

5% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than $2\mu\text{m}$ as measured by the Sedigraph 5100

0.1% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrilphenol methacrylate with 25 moles of ethylene oxide

and placed in aqueous suspension at a 59.8% concentration of dry substance using water and 0.67% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda and having a specific viscosity of 0.54.

Test N° 25 :

90% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 62% of the particles are of a diameter of less than 1 μ m as measured with the Sedigraph 5100

10% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μ m as measured by the Sedigraph 5100

0.2% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension at a 59.8% concentration of dry substance using water and 0.63% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda and having a specific viscosity of 0.54.

Test N° 26 :

85% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μ m as measured with the Sedigraph 5100

15% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μ m as measured by the Sedigraph 5100

0.3% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension at a 34.0% concentration of dry substance using water and 0.78% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate having a specific viscosity of 0.54.

Test N° 27 :

80% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 62% of the particles are of a diameter of less than 1 μm as measured with the Sedigraph 5100

20% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured by the Sedigraph 5100

0.4% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension at a 59.7% concentration of dry substance using water and 0.56% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda and having a specific viscosity of 0.54.

Test N° 28 :

70% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than $1\mu\text{m}$ as measured with the Sedigraph 5100

30% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than $2\mu\text{m}$ as measured by the Sedigraph 5100

0.6% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrilphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension with a 37.5% concentration of dry substance using water and 0.64% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate having a specific viscosity of 0.54.

Test N° 29 :

70% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 62% of the particles are of a diameter of less than $1\mu\text{m}$ as measured with the Sedigraph 5100

30% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than $2\mu\text{m}$ as measured by the Sedigraph 5100

0.6% by dry weight, relative to the total dry weight
 of fillers, of an acrylic copolymer binding
 agent comprising 90% by weight of acrylic acid
 and 10% by weight of tristyrylphenol
 methacrylate with 25 moles of ethylene oxide
 and placed in aqueous suspension at a 58.0%
 concentration of dry substance using water and 0.49%
 by dry weight, relative to the total dry weight of
 fillers, of a partially neutralised sodium
 polyacrylate having a specific viscosity of 0.5.

Test N° 30 :

This test, illustrating the prior art, is a simple
 mixture of 700 grams dry weight of an aqueous suspension
 of a Norwegian marble with a grain size equivalent to one
 in which 62% of the particles are of a diameter of less
 than 1 μm as measured on the Sedigraph 5100 and
 containing 1% by dry weight of a sodium polyacrylate with
 a specific viscosity of 0.7 with 300 grams dry weight of
 an aqueous suspension of Finnish talc with a grain size
 equivalent to one in which 45% of the particles are of a
 diameter of less than 2 μm as measured by the Sedigraph
 5100 containing 0.08% by dry weight of soda, 1.4% by dry
 weight of an alkylene polyoxide and 0.15% by dry weight
 of a sodium polyacrylate with a specific viscosity of 0.4
 to produce an aqueous suspension with a 66.4%
 concentration of a marble-talc mixture.

Test N° 31 :

For this test, illustrating the invention, the following
 composite compound proposed by the invention is prepared,
 using the same equipment and the same manner as test
 N° 29.

50% by dry weight, relative to the total dry weight

5 50% by dry weight, relative to the total dry weight
 of fillers, of Finnish talc with a grain size
 equivalent to one in which 45% of the particles
 are of a diameter of less than 2 μm as measured
 by the Sedigraph 5100

15 and placed in aqueous suspension at a 59.8%
concentration of dry substance using water and 0.7%
by dry weight, relative to the total dry weight of
fillers, of a partially neutralised sodium
polyacrylate having a specific viscosity of 0.5 and
20 0.2% by dry weight, relative to the total dry weight
of fillers, of a naphthalene sulphonic acid
condensate.

25 As in the previous test, the following composite compound
according to the invention is prepared.

25% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 62% of the particles are of a diameter of less than 1 μm as measured with the Sedigraph 5100

75% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles

are of a diameter of less than 2 μm as measured by the Sedigraph 5100

1.5% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension at a 56.6% concentration of dry substance using water and 0.63% by dry weight, relative to the total dry weight of fillers, of a partially neutralised sodium polyacrylate having a specific viscosity of 0.5 and 0.05% by dry weight, relative to the total dry weight of fillers, of a naphthalene-sulphonate acid condensate.

EXAMPLE 3 :

This example illustrates the preparation of composite compounds of the invention using different quantities of binding agent for a same pigment or filler composition.

To this end, the composite compounds proposed by the invention are prepared using the same method and the same equipment as in example 2 and comprise:

- 75% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 62% of the particles are of a diameter of less than 1 μm as measured with the Sedigraph 5100
- 25% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured by the Sedigraph 5100

and, as a binder, a same binding agent in different quantities.

These different quantities of binding agent tested,
5 acrylic copolymer of a monomer composition comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide, are as follows:

10 Test N°33 :

0.13% of binding agent by dry weight relative to the total dry weight of fillers.

15 An aqueous suspension with a 36.8% concentration of dry substances is formed using water and 0.69% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate with a specific viscosity of 0.54.

Test N°34 :

20 0.25% of binding agent by dry weight relative to the total dry weight of fillers.

25 An aqueous suspension with a 36.6% concentration of dry substances is formed using water and 0.69% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate with a specific viscosity of 0.54.

Test N°35 :

30 0.38% of binding agent by dry weight relative to the total dry weight of fillers.

An aqueous suspension with a 36.7% concentration of dry substances is formed using water and 0.69% by dry weight, relative to the total dry weight of fillers, of a sodium

polyacrylate with a specific viscosity of 0.54.

Test N° 36 :

1.25% of binding agent by dry weight relative to the
5 total dry weight of fillers.

An aqueous suspension with a 36.1% concentration of dry
substances is formed using water and 0.69% by dry weight,
relative to the total dry weight of fillers, of a sodium
10 polyacrylate with a specific viscosity of 0.54.

EXAMPLE 4 :

This example illustrates the preparation of composite
compounds as proposed by the invention using fillers or
15 pigments of different grain sizes.

To this end, the composite compounds proposed by the
invention are prepared by the same method and using the
same equipment as in example 2 and comprise:
20

Test N° 37 :

75% by dry weight, relative to the total dry weight
of fillers, of Norwegian marble with a grain
size equivalent to one in which 62% of the
25 particles are of a diameter of less than 1 μ m
as measured with the Sedigraph 5100

25% by dry weight, relative to the total dry weight
of fillers, of Finnish talc with a grain size
equivalent to one in which 45% of the particles
30 are of a diameter of less than 2 μ m as measured
by the Sedigraph 5100

0.5% by dry weight, relative to the total dry weight
of fillers, of an acrylic copolymer binding
agent comprising 90% by weight of acrylic acid

and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension at a 36.5% concentration of dry substance using water and 0.69% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate having a specific viscosity of 0.54.

Test N° 38 :

75% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 35% of the particles are of a diameter of less than 1 μm as measured with the Sedigraph 5100

25% by dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μm as measured by the Sedigraph 5100

0.5% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide and placed in aqueous suspension at a 36.4% concentration of dry substance using water and 0.69% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate having a specific viscosity of 0.54.

Test N° 39 :

75% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 75% of the

particles are of a diameter of less than 1 μm
as measured with the Sedigraph 5100

25% by dry weight, relative to the total dry weight
of fillers, of Australian talc with a grain
size equivalent to one in which 25% of the
particles are of a diameter of less than 2 μm
as measured by the Sedigraph 5100

0.5% by dry weight, relative to the total dry weight
of fillers, of an acrylic copolymer binding
agent comprising 90% by weight of acrylic acid
and 10% by weight of tristyrilphenol
methacrylate with 25 moles of ethylene oxide
and placed in aqueous suspension at a 35.4%
concentration of dry substance using water and 0.52%
by dry weight, relative to the total dry weight of
fillers, of a polyacrylate partially neutralised by
soda, having a specific viscosity of 0.5.

Test N° 40 :

75% by dry weight, relative to the total dry weight
of fillers, of Norwegian marble with a grain
size equivalent to one in which 75% of the
particles are of a diameter of less than 1 μm
as measured with the Sedigraph 5100

25% by dry weight, relative to the total dry weight
of fillers, of American talc with a grain size
equivalent to one in which 35% of the particles
are of a diameter of less than 2 μm as measured
by the Sedigraph 5100

0.5% by dry weight, relative to the total dry weight
of fillers, of an acrylic copolymer binding
agent comprising 90% by weight of acrylic acid
and 10% by weight of tristyrilphenol
methacrylate with 25 moles of ethylene oxide

and placed in aqueous suspension at a 36.1% concentration of dry substance using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised by soda having a specific viscosity of 0.5.

Test N° 41 :

50% by dry weight, relative to the total dry weight of fillers, of Champagne chalk with a grain size equivalent to one in which 36% of the particles are of a diameter of less than 2 μm as measured with the Sedigraph 5100

50% by dry weight, relative to the total dry weight of fillers, of Australian talc with a grain size equivalent to one in which 25% of the particles are of a diameter of less than 2 μm as measured by the Sedigraph 5100

2% by dry weight, relative to the total dry weight of fillers, of an acrylic copolymer binding agent comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide

and placed in aqueous suspension at a 59% concentration of dry substance using water and 0.35% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised by soda having a specific viscosity of 0.5.

Test N° 42 :

This test is conducted as a comparison with the preceding test and illustrates the preparation of an aqueous suspension known from the prior art using a standard mixture of

50% by dry weight, relative to the total dry weight

of fillers, of Champagne chalk with a grain size equivalent to one in which 36% of the particles are of a diameter of less than 2 μ m as measured with the Sedigraph 5100 and containing 0.07% by dry weight of a sodium polyacrylate with a specific viscosity of 0.7
50% by dry weight, relative to the total dry weight of fillers, of an aqueous suspension of Australian talc with a grain size equivalent to one in which 25% of the particles are of a diameter of less than 2 μ m as measured by the Sedigraph 5100 containing 0.08% by dry weight of soda, 1.4% by dry weight of an alkylene polyoxide and 0.15% by dry weight of a sodium polyacrylate with a specific viscosity of 0.4
to produce an aqueous suspension with a 71.7% concentration of dry substance of a chalk-talc mixture.

EXAMPLE 5 :

This example relates to the use of different binding agents.

To this end, the composite compounds proposed by the invention are made using the same method and the same equipment as in test N° 2, using as fillers 75% by dry weight, relative to the total dry weight of fillers, of Norwegian marble with a grain size equivalent to one in which 75% of the particles are of a diameter of less than 1 μ m as measured by the Sedigraph 5100 and 25% dry weight, relative to the total dry weight of fillers, of Finnish talc with a grain size equivalent to one in which 45% of the particles are of a diameter of less than 2 μ m as measured by the Sedigraph 5100 and, as a binder, different quantities of the following different binding agents:

Test N°43 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a polyacrylic acid with a specific viscosity of 1.78.

5

An aqueous suspension with a 59.7% concentration of dry substances is formed using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

10

Test N°44 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a polyacrylic acid with a specific viscosity of 1.55.

15

An aqueous suspension with a 60.4% concentration of dry substances is formed using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

20

Test N°45 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a polyacrylic acid with a specific viscosity of 0.95.

25

An aqueous suspension with a 59.8% concentration of dry substances is formed using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

30

Test N° 46 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is an acid polyacrylate, 10% neutralised by soda, having a specific viscosity of 5.00.

An aqueous suspension with a 59.9% concentration of dry substances is formed using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

Test N° 47 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a homopolymer of the methacrylate of ketostearyllic alcohol.

An aqueous suspension with a 59.2% concentration of dry substances is formed using water and 0.45% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

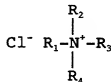
Test N° 48 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a copolymer comprising 98% by weight of methacrylic acid and 2% by weight of ketostearyllic alcohol methacrylate.

An aqueous suspension with a 59.7% concentration of dry substances is formed using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

Test N° 49 :

0.025% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a quaternary ammonium chloride having the formula



where R_1 = methyl radical

$\text{R}_2 = \text{R}_3$ = lauryl radical

R_4 = benzyl radical.

An aqueous suspension with a 59.3% concentration of dry substances is formed using water and 0.52% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

Test N° 50 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a linear alcohol having 12 carbon atoms.

An aqueous suspension with a 55.0% concentration of dry substances is formed using water and 0.75% by dry weight, relative to the total dry weight of fillers, of a polyacrylate partially neutralised with soda having a specific viscosity of 0.5.

Test N° 51 :

0.5% by dry weight, relative to the total dry weight of fillers, of a binding agent which is a linear alcohol having 18 carbon atoms.

An aqueous suspension with a 55.1% concentration of dry substances is formed using water and 0.38% by dry weight, relative to the total dry weight of fillers, of a sodium polyacrylate having a specific viscosity of 0.54.

Test N° 52 :

For this test, illustrating the invention, the aqueous co-structured compound proposed by the invention is prepared by introducing into a mixer under agitation 250 grams dry weight of a Finnish talc of a grain size such that 45% of the particles are of a diameter of less than 2 μ m as measured on the Sedigraph 5100, 5 grams dry weight of mixture of monomers comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide, 115 grams of isopropanol as well as the quantity of water needed to make up an aqueous composition with a 30% concentration of dry substance.

After 30 minutes of agitation, the mixture of monomers is polymerised, using known processes of radical polymerisation in a hydro-alcoholic medium.

Once the polymerisation is complete and the isopropanol has been removed by distillation, 750 grams dry weight of Norwegian marble are added, having a grain size such that 75% of the particles are of a diameter less than 1 μ m as measured by the Sedigraph 5100.

Once the co-structure has formed between the grains of talc and marble after 30 minutes of agitation, 7 grams by dry weight of a dispersing agent, namely a polyacrylic acid with a specific viscosity of 0.53, are added to obtain an aqueous suspension of the composite compound of

the invention having a concentration of dry substance equal to 36.4%.

Test N° 53 :

5 This test, illustrating the invention, using the same method and the same equipment as the preceding test, is conducted with 250 grams dry weight of a Finnish talc of a grain size such that 45% of the particles are of a diameter of less than 2 μm as measured on the Sedigraph
10 5100, 5 grams dry weight of a mixture of monomers comprising 90% by weight of acrylic acid and 10% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide, to which are added 5 grams by dry weight of a copolymer comprising 90% by dry weight of acrylic
15 acid and 10% by dry weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide, 115 grams of isopropanol as well as the quantity of water needed to make up an aqueous composition with a 30% concentration of dry substance.

20 After 30 minutes of agitation, the mixture of monomers is polymerised, using known processes of radical polymerisation in a hydroalcoholic medium.

25 Once the polymerisation is complete and the isopropanol has been removed by distillation, 750 grams dry weight of Norwegian marble are added, having a grain size such that 75% of the particles are of a diameter less than 1 μm as measured by the Sedigraph 5100.

30 Once the co-structure has formed between the grains of talc and marble after 30 minutes of agitation, 7 grams by dry weight of a dispersing agent, namely a polyacrylic acid with a specific viscosity of 0.53, are added to

obtain an aqueous suspension of the composite compound of the invention having a concentration of dry substance equal to 36.6%.

5 Test N° 54 :

This test, illustrating the invention, using the same method and the same equipment as the preceding test, is conducted with 250 grams dry weight of Finnish talc of a grain size such that 45% of the particles are of a
10 diameter of less than 2 μ m as measured on the Sedigraph 5100, 12.5 grams dry weight of a mixture of monomers comprising 80% by weight of acrylic acid and 20% by weight of tristyrylphenol methacrylate with 25 moles of ethylene oxide, 115 grams of isopropanol as well as the
15 quantity of water needed to make up an aqueous composition with a 30% concentration of dry substance.

After 30 minutes of agitation, the mixture of monomers is polymerised, using known processes of radical
20 polymerisation in a hydro-alcoholic medium.

Once the polymerisation is complete and the isopropanol has been removed by distillation, 750 grams dry weight of Norwegian marble are added, having a grain size such that
25 75% of the particles are of a diameter less than 1 μ m as measured by the Sedigraph 5100.

Once the co-structure has formed between the grains of talc and marble after 30 minutes of agitation, 7 grams by
30 dry weight of a dispersing agent, namely a polyacrylic acid with a specific viscosity of 0.53, are added to obtain an aqueous suspension of the composite compound of the invention having a concentration of dry substance equal to 36.6%.

Test N° 55 :

This test, illustrating the invention, using the same method and the same equipment as the preceding test, is conducted with 250 grams dry weight of a Finnish talc of a grain size such that 45% of the particles are of a diameter of less than 2 μ m as measured on the Sedigraph 5100, 5 grams dry weight of stearyl methacrylate, 115 grams of isopropanol as well as the quantity of water needed to make up an aqueous composition with a 30% concentration of dry substance.

After 30 minutes of agitation, the monomer is polymerised, using known processes of radical polymerisation in a hydro-alcoholic medium.

Once the polymerisation is complete and the isopropanol has been removed by distillation, 750 grams dry weight of Norwegian marble are added, having a grain size such that 75% of the particles are of a diameter less than 1 μ m as measured by the Sedigraph 5100.

Once the co-structure has formed between the grains of talc and marble after 30 minutes of agitation, 7 grams by dry weight of a dispersing agent, namely a polyacrylic acid with a specific viscosity of 0.53, are added to obtain an aqueous suspension of the composite compound of the invention having a concentration of dry substance equal to 36.7%.

EXAMPLE 6 :

This example is intended to demonstrate the formation of the co-structure or co-adsorption by measuring and comparing the homogeneity of the different suspensions of composite compounds obtained by diluting to a 20% concentration of dry substance.

To this end, the various aqueous suspensions of the composite compounds proposed by the invention and the suspensions of the prior art are diluted to a 20% concentration. Their macroscopic cohesion is measured using the homogeneity test consisting in determining the content of dry filler having at least one hydrophilic site at two separate points of the suspension representative of the test, namely a point located at the base of the flask and a point located on the surface of the flask after drying the suspension in an oven.

Once dry, the calcium ion content of each sample is determined after dissolving in HCl using EDTA sequestering techniques at a pH of 12 and with a coloured indicator, Eriochrome® Black T.

Test N° 56:

For this test, illustrating the prior art, the aqueous suspension of the mixture described in test N° 1 is used.

Test N° 57:

For this test, illustrating the invention, the aqueous suspension described in test N° 47 is used.

Test N° 58:

For this test, illustrating the invention, the aqueous suspension described in test N° 43 is used.

Test N° 59:

For this test, illustrating the invention, the aqueous suspension described in test N° 44 is used.

Test N° 60:

For this test, illustrating the invention, the aqueous

suspension described in test N°45 is used.

Test N°61:

For this test, illustrating the invention, the aqueous
5 suspension described in test N° 46 is used.

Test N°62:

For this test, illustrating the invention, the aqueous
10 suspension described in test N° 48 is used.

Test N°63:

For this test, illustrating the invention, the aqueous
suspension described in test N°49 is used.

15 Test N°64:

For this test, illustrating the invention, the aqueous
suspension described in test N° 50 is used.

Test N°65:

20 For this test, illustrating the invention, the aqueous
suspension described in test N°51 is used.

Test N°66:

For this test, illustrating the prior art, the aqueous
25 suspension described in test N°11 is used.

Test N° 67:

For this test, illustrating the invention, the aqueous
suspension of the composite compound of test N° 12 is
30 used.

Test N° 68:

For this test, illustrating the prior art, the aqueous suspension of the mixture described in test N° 13 is used.

5 Test N° 69:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 14 is used.

10 Test N° 70:

For this test, illustrating the prior art, the aqueous suspension described in test N° 15 is used.

15 Test N° 71:

For this test, illustrating the invention, the aqueous suspension described in test N° 16 is used.

20 Test N° 72:

For this test, illustrating the prior art, the aqueous suspension described in test N° 17 is used.

Test N° 73:

25 For this test, illustrating the invention, the aqueous suspension described in test N° 18 is used.

Test N° 74:

For this test, illustrating the prior art, the aqueous suspension described in test N° 19 is used.

30

Test N° 75:

For this test, illustrating the invention, the aqueous

suspension described in test N° 20 is used.

Test N° 76:

For this test, illustrating the prior art, the aqueous
5 suspension described in test N° 42 is used.

Test N° 77:

For this test, illustrating the invention, the aqueous
suspension described in test N° 41 is used.

10

Test N° 78:

For this test, illustrating the invention, the aqueous
suspension described in test N° 29 is used.

15 Test N° 79:

For this test, illustrating the prior art, the aqueous
suspension described in test N° 30 is used.

Test N° 80:

20 For this test, illustrating the invention, the aqueous
suspension described in test N° 52 is used.

Test N° 81:

For this test, illustrating the invention, the aqueous
25 suspension described in test N° 53 is used.

Test N° 82:

For this test, illustrating the invention, the aqueous
suspension described in test N° 54 is used.

30

Test N° 83:

For this test, illustrating the invention, the aqueous

suspension described in test N°55 is used.

Test N° 84:

For this test, illustrating the prior art, the aqueous suspension described in test N°23 is used.

It should be pointed out that in this test, the method used to measure the macroscopic homogeneity of the suspension is different from that of the preceding tests.

In effect, the quantities are not metered by sequestering techniques but by RFA analysis, which involves taking 0.2 g of the dry sample which is mixed with 1.625 g of lithium tetraborate, which is heated to melting temperature to obtain a disc, which is placed in a XRF 9400 machine made by ARL (Switzerland) to determine the elements by readings of the oxides present and then calculating the kaolin present.

The results of all the experiments are set out in Table 1 below.

- TABLE 1 -

	TEST N°	HOMOGENEITY % CaCO ₃ at the surface % CaCO ₃ at the base
Prior art	56	93.4 - 14.9
Invention	57	72.5 - 74.8
Invention	58	73.0 - 75.5
Invention	59	73.7 - 73.7
Invention	60	73.4 - 73.9
Invention	61	73.9 - 73.2
Invention	62	74.9 - 76.6
Invention	63	75.7 - 75.2
Invention	64	75.3 - 74.5
Invention	65	73.9 - 73.8
Prior art	66	74.7 - 63.9
Invention	67	74.4 - 73.4
Prior art	68	88.9 - 25.4
Invention	69	73.2 - 72.5
Prior art	70	90.2 - 37.1
Invention	71	83.1 - 83.5
Prior art	72	45.2 - 89.1
Invention	73	85.0 - 82.1
Prior art	74	29.6 - 85.5
Invention	75	81.7 - 80.5
Prior art	76	33.6 - 54.6
Invention	77	49.6 - 49.6
Invention	78	68.8 - 69.2
Prior art	79	91.9 - 32.5
Invention	80	74.8 - 74.1
Invention	81	74.6 - 73.6
Invention	82	74.4 - 75.4
Invention	83	70.3 - 72.7
Prior art	84	23.0 - 2.0*

5 * % kaolin at the surface - % kaolin at the base

Table I demonstrates that the aqueous suspensions containing the co-structured composite compounds of the invention have a more homogeneous content of fillers

with at least one hydrophilic site at different points than those containing the standard mixtures of the prior art.

5 EXAMPLE 7 :

This example demonstrates the formation of the co-structure or co-adsorption by measuring and comparing the viscosity and homogeneity of different paper coating colors produced.

10

To this end, the coating colors are prepared (tests 85 to 94) by mixing in water the composite compounds of fillers or pigments to be tested with

100 parts of compound to be tested at 65% of dry substance

12.5 parts of a carboxylated styrene-butadiene latex sold under the name of DL950 by Dow Chemical

and a quantity of water needed to obtain a 40% content of dry substances for tests N°85 to 92 and a content of dry substance in the order of 20% for tests N° 93 and 94.

25

The Brookfield viscosities of the coating colors prepared in this manner are then measured at ambient temperature at 20 revolutions/minute and 100 revolutions per minute using a Brookfield viscometer of the DVII type fitted with an appropriate spindle.

30

They are then tested for homogeneity using the same operating method as that described for the preceding example.

Test N°85:

For this test, illustrating a coating color of the

invention, the aqueous suspension of the composite compound of test N° 2 is used.

Test N° 86:

- 5 For this test, illustrating a coating color of the prior art, the aqueous suspension of the mixture of test N° 1 is used.

Test N° 87:

- 10 For this test, illustrating a coating color of the invention, the aqueous suspension of the composite compound of test N° 4 is used.

Test N° 88:

- 15 For this test, illustrating a coating color of the prior art, the aqueous suspension of the mixture of test N° 3 is used.

Test N° 89:

- 20 For this test, illustrating a coating color of the invention, the aqueous suspension of the composite compound of test N° 6 is used.

Test N° 90:

- 25 For this test, illustrating a coating color of the prior art, the aqueous suspension of the mixture of test N° 5 is used.

Test N° 91:

- 30 For this test, illustrating a coating color of the invention, the aqueous suspension of the composite compound of test N° 8 is used.

Test N° 92:

For this test, illustrating a coating color of the prior art, the aqueous suspension of the mixture of test N° 7 is used.

5

Test N° 93:

For this test, illustrating a coating color of the invention, the aqueous suspension of the composite compound of test N° 10 is used.

10

Test N° 94:

For this test, illustrating a coating color of the prior art, the aqueous suspension of the mixture of test N° 9 is used.

15

The results of all the experiments are set out in Table 2, the consistency of the coating colors of each of the tests being determined by introducing a spatula containing said coating colors.

20

- TABLE 2 -

	TES N°	VISCOSITY in mPa.s (20 r/mn)	VISCOSITY in mPa.s (100 r/mn)	CONSISTENCY	HOMOGENEITY % CaCO ₃ at surface % CaCO ₃ at the base
Invention	85	190	66	Soft	63.5 - 63.1
Prior art	86	14	24	Hard	76.1 - 34.8
Invention	87	765	180	Soft	62.6 - 63.0
Prior art	88	110	60	Medium hard	75.5 - 22.3
Invention	89	75	50	Soft	61.2 - 64.1
Prior art	90	16	29	Hard	65.8 - 48.5
Invention	91	242	88	Soft	64.1 - 64.4
Prior art	92	18	20	Hard	68.0 - 23.0
Invention	93	885	217	Soft	62.6 - 63.3
Prior art	94	55	47	Medium hard	66.4 - 50.5

Table 2 demonstrates that the coating colors of the invention containing the aqueous suspensions of co-structured composite compounds proposed by the invention are soft in appearance and have a higher Brookfield viscosity than that of the standard comparative mixtures of the prior art, thereby illustrating the co-structuring of the fillers or pigments. It is also clear that they have a more homogeneous content of fillers having at least one hydrophilic site at different points of the coating colors than that contained in the standard mixtures of the prior art.

EXAMPLE 8 :

This example relates to the measurements taken on the rheological behaviour of different aqueous compounds prepared in accordance with the method used in example 1.

The rheological behaviour of the different aqueous suspensions prepared in the same way as that used for example 1 is measured for viscoelasticity at 20°C using a Stress Tech® device by Reologica Instruments AB (Sweden) fitted with CC25 coaxial cylinders.

The same method was used to measure the rheological behaviour of the suspension produced in each of the tests, namely a sample of the suspension to be tested is injected into the cylinder of the viscoelasticity-measuring device and is pre-stressed at 10 Pa for 12 seconds and, after a waiting time of 180 seconds, stress is applied at a linear progression from 0.025 Pa to 20 Pa in 100 seconds and 40 intervals.

The yield stress, corresponding to the stress applied to the suspension to break the internal bonds and obtain a suspension of a reduced viscosity is determined by the maximum value of the viscosity curve in Pa.s as a function of the stress in Pa.

Test N°95:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 2 is used.

Test N°96:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°1 is used.

Test N°97:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 4 is used.

Test N°98:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°3 is used.

Test N°99:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 6 is used.

Test N°100:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°5 is used.

Test N° 101:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 8 is used.

Test N° 102:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N° 7 is used.

Test N° 103:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 10 is used.

Test N° 104:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N° 9 is used.

Test N° 105:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 12 is used.

Test N° 106:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N° 11 is used.

Test N° 107:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 14 is used.

Test N°108:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°13 is used.

5 Test N°109:

For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°15 is used.

Test N°110:

- 10 For this test, illustrating the invention, the aqueous suspension of the composite compound of test N°16 is used.

Test N°111:

- 15 For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°17 is used.

Test N°112:

- 20 For this test, illustrating the invention, the aqueous suspension of the composite compound of test N°18 is used.

Test N°113:

- 25 For this test, illustrating the invention, the aqueous suspension of the composite compound of test N°41 is used.

Test N°114:

- 30 For this test, illustrating the prior art, the aqueous suspension of the mixture of test N°42 is used.

Test N° 115:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 52 is used.

Test N° 116:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 53 is used.

Test N° 117:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 54 is used.

Test N° 118:

For this test, illustrating the invention, the aqueous suspension of the composite compound of test N° 55 is used.

The results of all the experiments are set out in Table 3 below:

- TABLE 3 -

	TEST N°	VISCOSITY in Pa.s	YIELD STRESS in Pa
Invention	95	642	3.072
Prior art	96	6.90	0.04465
Invention	97	164	0.9573
Prior art	98	1.49	0.03728
Invention	99	14 700	8.141
Prior art	100	0.527	0.03056
Invention	101	235	0.5842
Prior art	102	3.07	0.02965
Invention	103	1330	1.708
Prior art	104	38.4	0.3594
Invention	105	286	0.709
Prior art	106	12.32	0.079
Invention	107	2157	4.824
Prior art	108	4.81	0.102
Prior art	109	1.56	0.047
Invention	110	92	0.445
Prior art	111	43.2	0.099
Invention	112	589	0.336
Invention	113	938	1.580
Prior art	114	40.6	0.185
Invention	115	222	0.395
Invention	116	8.6	0.149
Invention	117	339	0.741
Invention	118	9.1	0.198

Table 3 demonstrates that the aqueous suspensions of the co-structured composite compound of the invention have a higher yield stress than that of the standard comparative mixtures of the prior art, characteristic of suspensions exhibiting good stability.

EXAMPLE 9 :

This example demonstrates the quality of the printability of the different paper coating colors produced in example 7.

This printability test, known as ISIT (Ink Surface Interaction Test) is based on a printing machine fitted with a device for creating and measuring the force needed to separate a contact disk from a film of printing ink. This machine consists on the one hand of the device which creates and measures the force and on the other an inking disk which rotates above the sheet of paper to be tested and is sold under the name of « Ink Surface Interaction Tester » by SeGan Ltd.

To this end, the different sheets of paper to be tested are prepared by applying the various coating colors to be tested on these sheets of paper using the Erichsen Model 624 laboratory coater made by Erichsen GmbH & Co. KG (Germany) fitted with detachable rotating blade coaters.

Once it has been coated with 7.5 g/m^2 , the paper to be tested is fixed on a roll fitted with a two-sided adhesive tape. An offset ink is applied by placing the inking disk in contact with a width of 25 mm as it is rotated 180° . The printing speed and pressure are adjustable and are in the order of 0.5 m/s and 50 kg

respectively. The volume of ink is under standard conditions of 0.3 cm^3 thus applying a thickness of approximately 1 g/m^2 to the sheet of paper to be tested.

- 5 The printing process is followed by a series of measurements to determine the tack force repeated at pre-selected time intervals depending on the time taken to separate the tack disk (of the same dimensions as the printing disk) from the film of ink.

10

A coating of offset-print quality nitrile rubber is usually used for the tack disk but any equivalent material may be used.

- 15 The contact force between the tack disk and the ink is measured by a system which generates an electromagnetic force. The amplitude and duration of the contact force are adjusted until a uniform adhesion is reached between the surface of the film and the tack disk at the end of 3 seconds. Rotating the sheet of paper slightly whilst the electromagnetic force is applied will ensure a close contact and continuity of the film of ink. Once the magnetic force is stopped, the tack disk is pulled back from the printed film by the force of a tensed spring, this force being sufficient to separate the disk from the film of ink. A strain gauge, mounted between the tack disk and the spring, generates a signal which is recorded as the tack force.

- 25
30 The sequence is automatically repeated for 13 cycles.

On the first and the thirteenth cycles, the printing densities are measured using a Gretag D 186 densitometer.

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This operating method is used for each of the coating colors to be tested, as follows:

Test N°119:

- 5 Illustrating the invention, the coating color of test N°85 is used.

Test N°120:

- 10 Illustrating the prior art, the coating color of test N°86 is used.

Test N°121:

- 15 Illustrating the invention, the coating color of test N°87 is used.

Test N°122:

- 20 Illustrating the prior art, the coating color of test N°88 is used.

Test N°123:

- 25 Illustrating the invention, the coating color of test N°89 is used.

Test N°124:

- 30 Illustrating the prior art, the coating color of test N°90 is used.

Test N°125:

- 35 Illustrating the invention, the coating color of test N°91 is used.

Test N° 126:

Illustrating the prior art, the coating color of test N° 92 is used.

5 Test N° 127:

Illustrating the invention, the coating color of test N° 93 is used.

Test N° 128:

10 Illustrating the prior art, the coating color of test N° 94 is used.

The results of all the experiments are set out in Tables 4 and 5 below and graphs 1 to 5 appended to this application.

Table 4 contains the results showing values of the tack force as a function of time whilst Table 5 gives the values of the printing density for tests 119 to 122.

Graphs 1 to 5 show the force which has to be applied to separate the disk from the film after printing as a function of time and can be interpreted taking the following three phases into account:

(i) the rise time, which essentially relates to the absorption and penetration speed of the ink as it makes initial contact with the surface to be printed.

The microporosity and wettability of this surface are major factors in terms of the rise time of this force.

The longer the rise time to the maximum value of the force, the more efficiently the binder of the

ink is absorbed, the less the film of ink will be susceptible to splitting and the better the adhesion between the ink and the paper, thus producing a better result.

- 5
- (ii) The maximum value of the tack force, which measures on the one hand the adhesion of the layer of ink which is immobilised on contact, the printing substrate and, on the other hand, the cohesion with the ink contained on the surface of the substrate. Consequently, the higher this maximum value of tack force for a constant cohesion, the better the adhesion will be and the better the print will be.
- 10

- 15
- (iii) The fall time of the force, which represents the drying of the ink. The slower this fall time, the slower the ink will dry, the less likelihood there will be of splitting in the structure of the ink and the better the print will be.
- 20

- TABLE 4 -

	Test 119 Invention	Test 120 Prior art	Test 121 Invention	Test 122 Prior art	Test 123 Invention	Test 124 Prior art	Test 125 Invention	Test 126 Prior art	Test 127 Invention	Test 128 Prior art
Time 5 seconds	Force: 1.89	Force: 3.93	Force: 2.73	Force: 3.74	Force: 3.94	Force: 4.32	Force: 2.82	Force: 2.70	Force: 3.29	Force: 4.67
Time 15 seconds	Force: 5.22	Force: 6.39	Force: 5.52	Force: 5.98	Force: 6.61	Force: 5.34	Force: 5.08	Force: 4.99	Force: 6.60	Force: 6.55
Time 26 seconds	Force: 6.71	Force: 6.11	Force: 6.46	Force: 5.89	Force: 6.77	Force: 5.27	Force: 6.30	Force: 5.39	Force: 7.38	Force: 6.08
Time 37 seconds	Force: 7.19	Force: 5.68	Force: 6.85	Force: 5.52	Force: 6.36	Force: 4.84	Force: 6.37	Force: 5.28	Force: 7.22	Force: 5.30
Time 57 seconds	Force: 7.21	Force: 5.45	Force: 6.84	Force: 5.50	Force: 5.63	Force: 4.43	Force: 6.10	Force: 4.63	Force: 6.19	Force: 3.74
Time 79 seconds	Force: 6.77	Force: 4.61	Force: 6.64	Force: 4.88	Force: 5.44	Force: 3.89	Force: 5.46	Force: 3.94	Force: 5.90	Force: 3.67
Time 98 seconds	Force: 6.06	Force: 4.41	Force: 6.37	Force: 4.71	Force: 4.56	Force: 2.95	Force: 5.05	Force: 2.73	Force: 4.00	Force: 2.10
Time 139 seconds	Force: 5.34	Force: 3.25	Force: 5.38	Force: 4.30	Force: 3.31	Force: 2.42	Force: 4.16	Force: 1.65	Force: 3.25	Force: 1.25
Time 179 seconds	Force: 5.07	Force: 2.27	Force: 4.66	Force: 3.04	Force: 2.59	Force: 2.09	Force: 3.15	Force: 0.93	Force: 1.73	Force: 0.79
Time 220 seconds	Force: 4.54	Force: 1.67	Force: 3.80	Force: 2.22	Force: 2.37	Force: 1.57	Force: 2.78	Force: 1.39	Force: 1.36	Force: 0.72
Time 281 seconds	Force: 3.67	Force: 1.02	Force: 3.22	Force: 1.42	Force: 2.03	Force: 0.99	Force: 2.24	Force: 1.32	Force: 1.16	Force: 0.64
Time 341 seconds	Force: 3.12	Force: 0.88	Force: 2.67	Force: 1.01	Force: 1.21	Force: 0.90	Force: 1.55	Force: 1.19	Force: 0.71	Force: 0.59
Time 402 seconds	Force: 3.23	Force: 0.92	Force: 2.43	Force: 0.86	Force: 1.30	Force: 0.89	Force: 1.56	Force: 1.07	Force: 0.72	Force: 0.66

- TABLE 5 -

	Test N°	Print density 1st cycle	Print density 13th cycle
Invention	119	1.58	1.05
Prior art	120	1.49	0.97
Invention	121	1.52	1.05
Prior art	122	1.48	0.94

Table 4 and graphs N° 1 to 5 demonstrate that the coating colors proposed by the invention have slower rise and fall times and higher tack force values, which means that they exhibit better printability in terms of adhesion, brightness and print yield.

Table 5 demonstrates that the coating colors of the invention have higher print densities than those of the comparative coating colors known from the prior art.

EXAMPLE 10 :

- 15 This example relates to the opacity measurement and more specifically to determining the light scattering coefficient S of the various coating colors.

20 The method used to determine the light coefficient S, well known to those skilled in the art, is as follows:

For each test, a sheet of paper containing no wood is coated with the coating color to be tested.

- 25 Before being coated and prior to each test, this sheet of paper, the dimensions of which are 10 cm x 6 cm and

with a specific weight of 75.5 g/m², is weighed and then irradiated with light of a wavelength equal to 457 nm on a black plate using an Elrepho 2000 spectrophotometer by Datacolor (Switzerland) to determine the base reflection index R_b.

Each of the coating colors to be tested is then applied to this pre-weighed sheet of paper with a laboratory coater fitted with detachable rotating coating blades sold as reference model 624 by Erichsen (Germany). Each sheet of paper coated in this manner at 7.5 g/m² is then irradiated with light of a wavelength equal to 457 nm by means of an Elrepho 2000 spectrophotometer by Datacolor (Switzerland) on a black plate to determine the reflection index R₀ and on a stack of sheets of paper which have not been coated in order to determine the reflection index R₁, where r is the reflection index of the stack of uncoated sheets of paper.

The reflection index R_{sc} of the single coat, on a black background, is then determined using the formula:

$$R_{sc} = \frac{R_1 \cdot R_b - R_0 \cdot r}{(R_1 - R_0) \cdot r R_0 + R_b - r}$$

and the transmittance T_{sc} of the coating

$$T_{sc}^2 = \frac{(R_0 - R_{sc})(1 - R_{sc} R_b)}{R_b}$$

to obtain a theoretic reflection value R_∞ for a coating of infinite thickness given by the formula:

$$\frac{1 - T_{sc}^2 + R_{sc}^2}{R_{sc}} = \frac{1 + R_{sc}^2}{R_{sc}}$$

5 The scattering coefficient S characteristic of the opacity can be calculated from this formula, knowing that, for a coating weight P,

$$10 \quad S.P. = \frac{1}{b} \coth^{-1} \frac{(1 - aR_{sc})}{bR_{sc}}$$

$$a = 0.5 \left(\frac{1}{R_{sc}} + R_{sc} \right)$$

$$15 \quad \text{where } b = 0.5 \left(\frac{1}{R_{sc}} + R_{sc} \right)$$

Test N°129:

For this test, illustrating the prior art, the coating color of test N°86 is used.

Test N°130:

For this test, illustrating the invention, the coating color of test N°85 is used.

Test N°131:

For this test, illustrating the prior art, the coating color of test N°88 is used.

Test N°132:

For this test, illustrating the invention, the coating color of test N°87 is used.

All the results are set out in Table 6 below:

- TABLE 6 -

	PRIOR ART	INVENTION	PRIOR ART	INVENTION
TEST N°	129	130	131	132
S in m ² /kg	143	157	104	136

Table 6 demonstrates that the coating colors of the invention have a higher light scattering coefficient S than the comparative coating colors of the prior art.

EXAMPLE 11 :

This example relates to the direct opacity measurement and the whiteness of the coating colors in compliance with the TAPPI T452 ISO 2470 standard.

For each test, a sheet of paper containing no wood and of dimensions 10 cm x 6 cm, having a specific weight of 75.5 g/m², is coated with the coating color to be tested using a laboratory coating machine with removable rotating blade coaters, marketed as Model 624 by Erichsen (Germany).

Each sheet of paper coated in this manner with 7.5 g/m² is then exposed to light of a wavelength of 457 nm using an Elrepho 2000 spectrophotometer by Data Color (Switzerland) to determine the opacity and whiteness.

This example also relates to the brightness measurement. The brightness measurement is taken on the

same coated sheets of paper as those used to take the direct opacity and whiteness measurements.

This method involves feeding the coated sheet of paper into the laboratory brightness meter, LGDL -05/2 (Lehmann Messtechnik AG, Switzerland) which measures the 75° TAPPI brightness according to Lehmann.

Test N°133:

For this test, illustrating the prior art, the coating color of test N°88 is used.

Test N°134:

For this test, illustrating the invention, the coating color of test N°87 is used.

Test N°135:

For this test, illustrating the prior art, the coating color of test N°92 is used.

Test N°136:

For this test, illustrating the invention, the coating color of test N°91 is used.

Test N°137:

For this test, illustrating the prior art, the coating color of test N°94 is used.

Test N°138:

For this test, illustrating the invention, the coating color of test N°93 is used.

The results of the opacity measurement experiments are set out in Table 7 below:

- TABLE 7 -

	PRIOR ART	INVENTION
TEST N°	133	134
OPACITY	91.1%	92.2%

The results of the whiteness measurement experiments are set out in Table 8 below:

- TABLE 8 -

	PRIOR ART	INVENTION
TEST N°	135	136
WHITENESS	84.8%	87.6%

The results of the brightness measurement experiments are set out in Table 9 below:

- TABLE 9 -

	PRIOR ART	INVENTION
TEST N°	137	138
BRIGHTNESS	41.4%	48.6%

Tables 7 to 9 demonstrate that the coating colors of

the invention have a higher opacity, whiteness and brightness than those produced by the comparative coating colors of the prior art.

EXAMPLE 12 :

This example relates to the opacity measurement conforming to the DIN 53146 standard and the whiteness of sheets of paper containing, in the mass, the uncoated filling compounds proposed by the invention and comparisons of these with those containing the standard suspensions of mixtures of the prior art.

To this end, sheets of paper are made from a cellulose pulp of SR 23 degree containing a woodless sulphate paste and fibres comprising 80% birch and 20% pine. 45g dry weight of this pulp are then diluted in 10 litres of water in the presence of about 15 g dry weight of the filler composition to be tested to produce experimentally a filler content of 20%. After 15 minutes of agitation and adding 0.06% by dry weight, relative to the dry weight of paper, a retention agent of the polyacrylamide type, a sheet is made with a grammage equal to 75 g/m² and filled to 20%. The device used to make the sheet is a Rapid-Köthen 20.12 MC model by Haage.

The sheets prepared in this manner are dried for 400 seconds at 92°C and in a vacuum of 940 mbar. The filler content is controlled by analysing the ashes.

The different values for opacity and whiteness are then determined using the same method as before.

The following tests were conducted.

Test N°139:

For this test, illustrating the prior art, the mixture of test N°1 is used.

Test N°140:

For this test, illustrating the invention, the composite compound of test N°2 is used.

Test N°141:

For this test, illustrating the prior art, the mixture of test N°3 is used.

Test N°142:

For this test, illustrating the invention, the composite compound of test N°4 is used.

The results of the whiteness measurement experiments are set out in Table 10 below:

- TABLE 10 -

	PRIOR ART	INVENTION
TEST N°	139	140
WHITENESS	86.9	87.7

The results of the opacity measurement experiments are set out in Table 11 below:

TABLE 11

	PRIOR ART	INVENTION
TEST N°	141	142
OPACITY	88.7	90.3

Tables 10 and 11 demonstrate that the sheets filled with the co-structured composite compounds of the invention have a higher opacity and whiteness than those filled with the standard mixtures of the prior art.

EXAMPLE 13 :

This example relates to the opacity measurement and more specifically to determining the light scattering coefficient S of an aqueous paint composition containing, essentially water and 100 parts of the composite compound to be tested with a 65 % dry content as well as 9.8 parts of a binder polymeric dispersion of the styrene-acrylic type.

This coefficient is measured by using the same way that used for example 10, except that the support to be coated is not a sheet of paper but an aluminum sheet.

The obtained results are completely the same as those obtained in the tests No. 129 and No. 130, and so demonstrate that the paint compositions according to the invention have a higher light scattering S coefficient than a paint composition of the prior art

and so an increased opacity.

From the results of the example 10, the skilled man in the art was expecting the results of the example 13.

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In fact, this S coefficient, as confirmed by the disclosed calculation in the example 10, is independent from the support (paper or metal or cement sheet) and depends only from the coating composition i.e. from the paper coating color or from the paint composition.

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- CLAIMS -

1/ A composite compound of mineral or organic fillers or pigments, characterised in that it contains:

a) at least two mineral or organic fillers or pigments, at least one of which has a surface with at least one hydrophilic site and the other at least has at least one organophilic site

b) at least one binding agent

and in that the mineral or organic fillers or pigments are co-structured or co-adsorbed.

2/ A composite compound of mineral or organic fillers or pigments as claimed in claim 1, characterised in that it is an aqueous compound.

3/ A composite compound of mineral or organic fillers or pigments as claimed in claim 1, characterised in that it is a non-aqueous compound.

4/ A composite compound of mineral or organic fillers or pigments as claimed in claim 1, characterised in that it is a dry compound.

5/ A composite compound as claimed in one of claims 1 to 4, characterised in that the binding agent is an organic compound.

6/ A composite compound as claimed in any one of claims 1 to 5, characterised in that the binding agent is supported by a gas.

7/ A composite compound as claimed in any one of claims 1 to 6, characterised in that the binding agent is selected from among the acrylic or vinyl polymers and/or copolymers or polycondensates or the
5 polyaddition products such as polymers and/or copolymers, in their totally acid state or partially neutralised or totally neutralised, of at least one of the monomers such as acrylic and/or methacrylic, itaconic, crotonic, fumaric acid, maleic anhydride or
10 isocrotonic, aconitic, mesaconic, sinapic, undecylenic, angelic acid and/or their respective esters, acrylamido methyl propane sulphononic acid, acrolein, acrylamide and/or methacrylamide, methacrylamido propyl trimethyl ammonium chloride or sulphate, methacrylate of
15 trimethyl ammonium ethyl chloride or sulphate, as well as their acrylate and acrylamide counterparts, quaternised or not, and/or dimethyldiallyl chloride, vinylpyrrolidone or a binding agent chosen from among the linear or branched fatty acids or the linear or
20 branched fatty alcohols or the linear or branched or cyclic fatty amines, saturated or unsaturated, or a binding agent chosen from among the quaternary salts, preferably with linear or branched fatty chains of vegetable origin or not.

8/ A composite compound as claimed in any one of claims 1 to 6, characterised in that the binding agent is selected from among the acrylic or vinyl polymers and/or copolymers. in their totally acid state or
30 partially neutralised or totally neutralised, obtained by polymerisation, in the acid state in the presence of one at least of the mineral or organic particles of the composite compound and optionally in the presence of the binding agent as claimed in claim 7, of at least
35 one of the monomers such as acrylic and/or methacrylic,

itaconic, crotonic, fumaric acid, maleic anhydride or isocrotonic, aconitic, mesaconic, sinapic, undecylenic, angelic acid and/or their respective esters, acrylamido methyl propane sulphonc acid, acrolein, acrylamide and/or methacrylamide, methacrylamido propyl trimethyl ammonium chloride or sulphate, methacrylate of trimethyl ammonium ethyl chloride or sulphate, as well as their acrylate and acrylamide counterparts, quaternised or not, and/or dimethyldiallyl chloride, vinylpyrrolidone or such as the unsaturated linear or branched fatty acids or the linear or branched fatty alcohols or the unsaturated linear or branched or cyclic fatty amines, or the quaternary salts, preferably with unsaturated linear or branched fatty chains of vegetable origin or not.

9/ A composite compound as claimed in any one of claims 1 to 8, characterised in that the mineral or organic filler or fillers and pigment or pigments having a surface with at least one hydrophilic site are chosen from among the natural calcium carbonates such as chalk, calcite, marble or any other form of natural calcium carbonate, obtained in particular from recycling processes, or from among precipitated calcium carbonate, the dolomites, crystalline or amorphous aluminium hydroxides, synthetic or natural precipitated silicates, calcium sulphate, titanium dioxides, satin white, the wollastonites, huntite, calcined clays in particular from recycling processes or starch or alternatively selected from among organophilic organic or mineral particles that have been subjected to physical or chemical processing so that they have at least one hydrophilic site.

10/ A composite compound as claimed in any one of claims 1 to 8, characterised in that the mineral or organic filler or fillers and pigment or pigments having a surface with at least one organophilic site are selected from among the talcs, micas, clays, whether calcined or not, zinc oxide, transparent iron pigments, colouring pigments, synthetic pigments with a polystyrene base, urea-formol resins, carbon black or the fibres and flour of cellulose or selected from among hydrophilic mineral or organic particles that have been subjected to physical or chemical processing so that they have at least one organophilic site.

11/ A composite compound as claimed in any one of claims 1 to 10, characterised in that it contains 0.1% to 99.9% by dry weight and preferably 25% to 95% by dry weight, relative to the total dry weight of the fillers or pigments, of mineral or organic fillers or pigments having a surface with at least one hydrophilic site and 99.9% to 0.1% by dry weight and preferably between 75% and 5% by dry weight, relative to the total dry weight of the fillers or pigments, of mineral or organic fillers or pigments having a surface with at least one organophilic site.

12/ A composite compound as claimed in any one of claims 1 to 11, characterised in that it contains 0.01% to 10% and preferably 0.1% to 1.5% dry weight of the binding agent relative to the total dry weight of the fillers or pigments.

13/ A composite compound as claimed in any one of claims 1 to 12, characterised in that it is macroscopically homogeneous.

14/ A composite compound as claimed in any one of claims 1 to 12, characterised in that its yield stress as determined by measuring on a Stress Tech® machine for measuring viscoelasticity is higher than and preferably at least four times higher than that of the standard corresponding standard mixture of fillers or pigments.

15/ Use of the composite compounds as claimed in any one of claims 1 to 14 for the manufacture of aqueous suspensions of organic or mineral pigments or fillers, paper coating colors and/or for the manufacture of paper, in the paper filling and/or for any other paper surface-treatment.

16/ Use of the composite compounds as claimed in any one of claims 1 to 14 in the field of paints.

17/ Use of the composite compounds as claimed in any one of claims 1 to 14 in the field of plastics.

18/ An aqueous suspension of mineral or organic fillers or pigments, characterised in that it contains a composite compound as claimed in any one of claims 1 to 14.

19/ An aqueous suspensions of mineral or organic fillers or pigments as claimed in claim 18, characterised in that it is macroscopically homogeneous.

20/ An aqueous suspension of mineral or organic fillers or pigments as claimed in claim 18, characterised in that its yield stress as determined by a Stress Tech®

machine for measuring viscoelasticity is higher than and preferably at least four times higher than that of the corresponding standard mixture of fillers or pigments.

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21/ A coating color characterised in that it contains a composite compound as claimed in any one of claims 1 to 14.

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22/ A paper coating color as claimed in claim 21 characterised in that it is macroscopically homogeneous.

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23/ A paper coating color as claimed in claim 21, characterised in that its yield stress as determined by a Stress Tech® machine for measuring viscoelasticity is higher than and preferably at least four times higher than the corresponding standard mixture of fillers or pigments.

20

24/ A paper coating color as claimed in any one of claims 21 to 23, characterised in that it has a higher light scattering coefficient S than that of a coating color containing the standard suspensions of the corresponding mixtures.

25

25/ A paper coating color as claimed in any one of claims 21 to 23, characterised in that it has a higher whiteness, determined in accordance with the TAPPI T452 ISO 2470 standard, than that of a coating color containing standard suspensions of the corresponding mixtures.

30

26/ A paper coating color as claimed in any one of

claims 21 to 23, characterised in that it has a higher brightness, TAPPI 75° according to Lehmann, than that of a coating color containing the standard suspensions of corresponding mixtures.

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27/ A paper coating color as claimed in any one of claims 21 to 23, characterised in that its curve, determined in accordance with the ISIT printability test and representative of the tack force as a function of time, has smaller rising and falling gradients than coating colors containing the standard suspensions of the corresponding mixtures and a higher maximum value in terms of tack force.

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28/ A paper coating color as claimed in any one of claims 21 to 23, characterised in that it has a higher print density than that of a coating color containing the standard suspensions of the corresponding mixtures.

20

29/ A paper surface-treatment compound as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous paint or non-aqueous compositions characterised in that it contains a composite compound as claimed in any one of claims 1 to 14.

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30/ A paper surface-treatment compound as well as wood or metal or plastic or cement surface treatment compounds and/or aqueous paint or non-aqueous compositions as claimed in claim 29, characterised in that it is macroscopically homogeneous.

35

31/ A paper surface-treatment compound as claimed in claim 29, characterised in that its yield stress as determined by the Stress Tech® machine for measuring

viscoelasticity is higher and preferably at least four times higher than that of the standard corresponding mixture of fillers or pigments.

- 5 32/ Aqueous or non-aqueous paint composition as claimed in claim 29, characterised in that it has a higher light scattering coefficient S than that of a paint composition containing the standard suspensions of the corresponding mixtures.

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33/ A paper surface-treatment compound as claimed in any one of claims 29 to 31, characterised in that its curve, determined in accordance with the ISIT printability test and representative of the tack force as a function of time, has smaller rising and falling gradients than surface-treatment compounds containing the standard suspensions of the corresponding mixtures and a higher maximum value in terms of tack force.

15

- 20 34/ An uncoated filling composition characterised in that it contains a composite compound as claimed in any one of claims 1 to 14.

35/ A sheet of base paper to be coated, characterised in that it contains the uncoated filling composition as claimed in claim 34.

25

36/ A sheet of paper as claimed in claim 35, characterised in that it has a higher opacity determined in accordance with the DIN 53146 standard than that of a sheet of paper containing the standard suspensions of corresponding mixtures.

30

37/ A sheet of paper as claimed in claim 35, characterised in that it has a higher whiteness

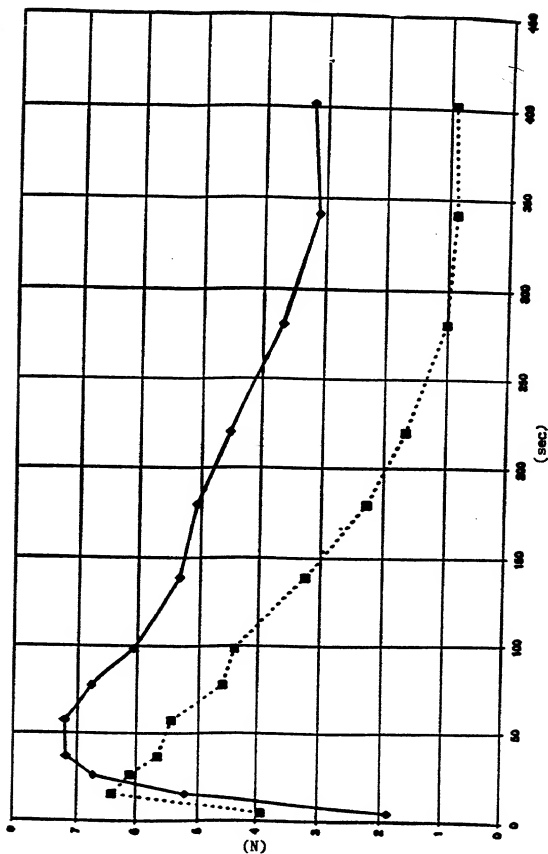
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09/646897

GRAPH 1

TEST N° 119

TEST N° 120



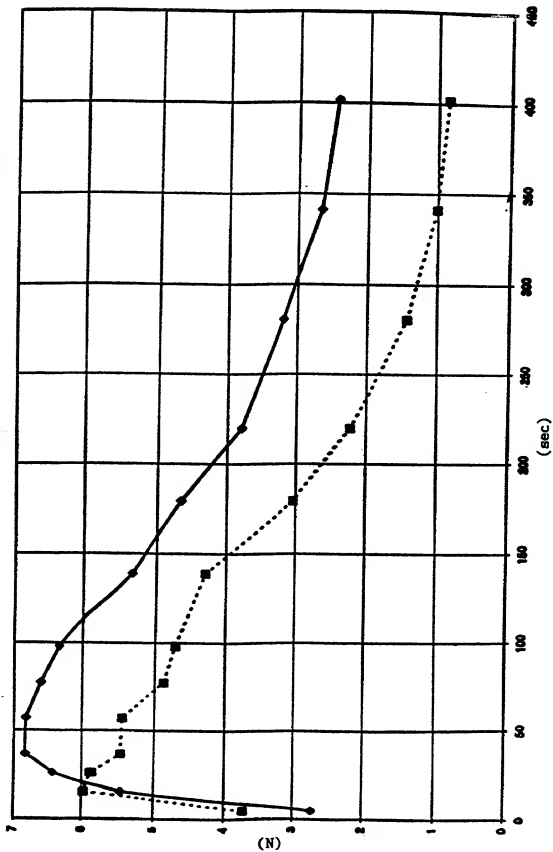
09/646897.121100

09/646897

GRAPH 2

TEST N° 121

TEST N° 122

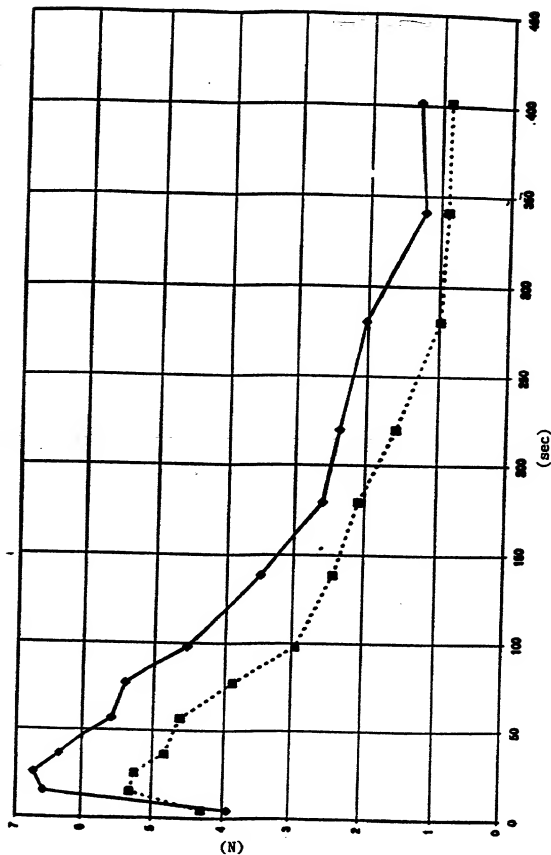


09646897.121100

09/646897

TEST N° 123

GRAPH 3 TEST N° 124



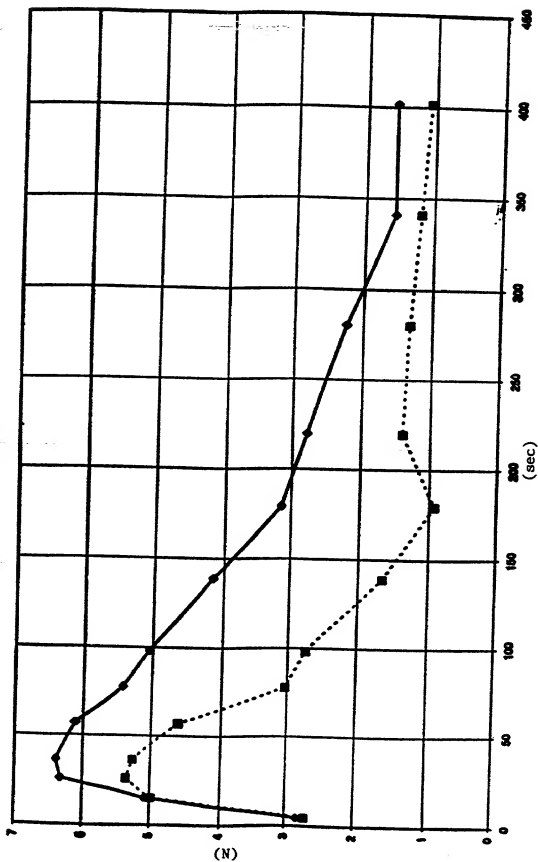
09646897.124100

09/646897

TEST N° 125

TEST N° 126

GRAPH 4



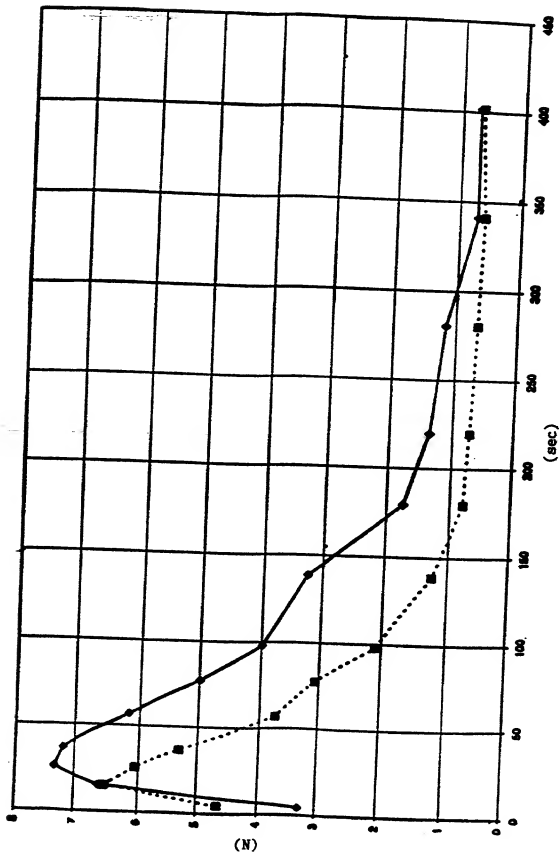
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TEST N° 127

GRAPH 5

TEST N° 128



Declaration, Power Of Attorney and Petition

Page 1 of 3

WE (I) the undersigned inventor(s), hereby declare(s) that:

My residence, post office address and citizenship are as stated below next to my name,

We (I) believe that we are (I am) the original, first, and joint (sole) inventor(s) of the subject matter which is claimed and for which a patent is sought on the invention entitled

COMPOSITE CO-STRUCTURED OR CO-ADSORBED, MINERAL OR ORGANIC FILLER OR PIGMENT
COMPOUNDS AND THE USE THEREOF ✓

the specification of which

☐ is attached hereto.

☒ was filed on October 10, 2000 ✓ as
Application Serial No. 09/646,897 ✓
and amended on _____.

☒ was filed as PCT international application
Number PCT/IB99/00941 ✓
on April 06, 1999 ✓,
and was amended under PCT Article 19
on _____ (if applicable).

We (I) hereby state that we (I) have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We (I) acknowledge the duty to disclose information known to be material to the patentability of this application as defined in Section 1.56 of Title 37 Code of Federal Regulations.

We (I) hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s)

Application No.	Country	Day/Month/Year	Priority Claimed
<u>98/04714</u> ✓	<u>FRANCE</u> ✓	<u>09 April 1998</u> ✓	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
<u>98/15244</u> ✓	<u>FRANCE</u> ✓	<u>30 November 1998</u> ✓	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
_____	_____	_____	<input type="checkbox"/> Yes <input type="checkbox"/> No
_____	_____	_____	<input type="checkbox"/> Yes <input type="checkbox"/> No

We (I) hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below.

_____ (Application Number)	_____ (Filing Date)
_____ (Application Number)	_____ (Filing Date)

We (I) hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

Application Serial No.	Filing Date	Status (pending, patented, abandoned)
PCT/IB99/00941 ✓	06 April 1999	
_____	_____	_____
_____	_____	_____

And we (I) hereby appoint: Norman F. Oblon, Reg. No. 24,618; Marvin J. Spivak, Reg. No. 24,913; C. Irvin McClelland, Reg. No. 21,124; Gregory J. Maier, Reg. No. 25,599; Arthur I. Neustadt, Reg. No. 24,854; Richard D. Kelly, Reg. No. 22,757; James D. Hamilton, Reg. No. 28,421; Eckhard H. Kuesters, Reg. No. 28,870; Robert T. Pous, Reg. No. 29,099; Charles L. Gholz, Reg. No. 26,395; William E. Beaumont, Reg. No. 30,996; Jean-Paul Lavalleye, Reg. No. 31,451; Stephen G. Baxter, Reg. No. 32,884; Richard L. Treanor, Reg. No. 36,379; Steven P. Weihrouch, Reg. No. 32,829; John T. Goolkasian, Reg. No. 26,142; Richard L. Chinn, Reg. No. 34,305; Steven E. Lipman, Reg. No. 30,011; Carl E. Schlier, Reg. No. 34,426; James J. Kulbaski, Reg. No. 34,648; Richard A. Neifeld, Reg. No. 35,299; J. Derek Mason, Reg. No. 35,270; Surinder Sachar, Reg. No. 34,423; Christina M. Gadiano, Reg. No. 37,628; Jeffrey B. McIntyre, Reg. No. 36,867; William T. Enos, Reg. No. 33,128; Michael E. McCabe, Jr., Reg. No. 37,182; Bradley D. Lytle, Reg. No. 40,073; and Michael R. Casey, Reg. No. 40,294; our (my) attorneys, with full powers of substitution and revocation, to prosecute this application and to transact all business in the Patent Office connected therewith; and we (I) hereby request that all correspondence regarding this application be sent to the firm of OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT, P.C., whose Post Office Address is: Fourth Floor, 1755 Jefferson Davis Highway, Arlington, Virginia 22202.

We (I) declare that all statements made herein of our (my) own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

1-00 Patrick A. GANE ✓
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2-00 Matthias BURI
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M. Buri
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NAME OF THIRD JOINT INVENTOR

Signature of Inventor

Date

NAME OF FOURTH JOINT INVENTOR

Signature of Inventor

Date

NAME OF FIFTH JOINT INVENTOR

Signature of Inventor

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Residence: _____

Citizen of: _____

Post Office Address: _____

Residence: _____

Citizen of: _____

Post Office Address: _____

Residence: _____

Citizen of: _____

Post Office Address: _____